

97965

STIC-Biotech/ChemLib

From: Fredman, Jeffrey
Sent: Wednesday, July 02, 2003 12:51 PM
To: STIC-Biotech/ChemLib
Cc: Schultz, James
Subject: FW: Rush Sequence search request 09/780,929

PLEASE RUSH.

I Approve.

Jeff Fredman

CRF

-----Original Message-----

From: Schultz, James
Sent: Wednesday, July 02, 2003 9:38 AM
To: Fredman, Jeffrey
Subject: Rush Sequence search request 09/780,929

Dear Jeff,

Would you please approve the rush sequence search below? This case has already been searched and is ready for allowance, but my SPE wants it searched one more way before we pass it out.

Thanks,

Doug Schultz

Dear STIC-biotech searchers,

Could you please run a length limited nucleotide sequence search on SEQ ID NOS 97 (15 nt long) and 98 (18 nt long) in the above entitled case, where the maximum size of the returned hit is no longer than 60 nucleotides? I need both sequences searched in the **interference** databases as well.

Thanks,

Doug Schultz

J. Douglas Schultz, Ph.D.
AU 1635 (Biotechnology)
Patent Examiner
United States Patent and Trademark Office
(703) 308-9355
(703) 746-3973 (fax)
Office: CM1 12E18
Mail: CM1 11E12

Searcher: _____
Phone: _____
Location: _____
Date Picked Up: _____
Searcher Prep/Review: _____
Clerical: _____
Online time: _____

TYPE OF SEARCH:
NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST (where applic.)
STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: _____
WWW/Internet: _____
Other (specify): _____

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:51:21 ; Search time 520.455 seconds

(without alignments)
206.249 Million cell updates/sec

Title: US-09-780-929-97

Perfect score: 15

Sequence: 1 agauaacgugaagau 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 8255821 seqs, 3578102051 residues

Total number of hits satisfying chosen parameters: 9359164

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

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14: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq2:*
15: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq3:*
16: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq4:*
17: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq:*
18: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq2:*
19: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq3:*
20: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	13.4	89.3	25	14	US-10-355-577-24855
C 2	13.4	89.3	25	18	US-60-427-836-243678
C 3	13	86.7	25	14	US-10-355-577-179244
C 4	13	86.7	25	18	US-60-427-836-118114
C 5	12.4	82.7	23	14	US-10-310-188-75880
C 6	12.4	82.7	25	10	US-09-660-222-72811
C 7	12.4	82.7	25	10	US-09-660-222-72812
C 8	12.4	82.7	25	11	US-09-953-570-48322
C 9	12.4	82.7	25	11	US-09-953-570-48329
C 10	12.4	82.7	25	11	US-09-954-445A-10200
C 11	12.4	82.7	25	11	US-09-954-445A-16612
C 12	12.4	82.7	25	11	US-09-954-445A-16620
C 13	12.4	82.7	25	11	US-09-954-445A-50048

14	12.4	82.7	25	14	US-10-355-577-40591	Sequence 40591, A
C 15	12.4	82.7	25	14	US-10-355-577-366476	Sequence 366476, A
C 16	12.4	82.7	25	14	US-10-355-577-485354	Sequence 485354, A
C 17	12.4	82.7	25	14	US-10-355-577-511942	Sequence 511942, A
C 18	12.4	82.7	25	14	US-10-355-577-735345	Sequence 735345, A
C 19	12.4	82.7	25	14	US-10-355-577-834196	Sequence 834196, A
C 20	12.4	82.7	25	14	US-10-355-577-907847	Sequence 907847, A
C 21	12.4	82.7	25	18	US-60-427-808-475949	Sequence 475949, A
C 22	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
C 23	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
C 24	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
C 25	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
C 26	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
C 27	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
C 28	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
C 29	12.4	82.7	25	19	US-60-469-545-95587	Sequence 95587, A
C 30	12.4	82.7	25	19	US-60-469-545-95587	Sequence 95587, A
C 31	12.4	82.7	25	19	US-60-469-545-95587	Sequence 95587, A
C 32	12.4	82.7	25	19	US-60-469-545-95587	Sequence 95587, A
C 33	12.4	82.7	25	20	US-60-469-545-95587	Sequence 95587, A
C 34	12.4	82.7	25	20	US-60-469-545-95587	Sequence 95587, A
C 35	12	80.0	22	14	US-10-310-188-49187	Sequence 49187, A
C 36	12	80.0	22	14	US-10-310-188-49187	Sequence 49187, A
C 37	12	80.0	25	11	US-09-953-570-45886	Sequence 45886, A
C 38	12	80.0	25	11	US-09-953-570-45886	Sequence 45886, A
C 39	12	80.0	25	14	US-10-098-263B-27903	Sequence 27903, A
C 40	12	80.0	25	14	US-10-355-577-53971	Sequence 53971, A
C 41	12	80.0	25	14	US-10-355-577-143487	Sequence 143487, A
C 42	12	80.0	25	14	US-10-355-577-518580	Sequence 518580, A
C 43	12	80.0	25	14	US-10-355-577-518580	Sequence 518580, A
C 44	12	80.0	25	14	US-10-355-577-518580	Sequence 518580, A
C 45	12	80.0	25	14	US-10-355-577-827041	Sequence 827041, A
					US-10-355-577-899636	Sequence 899636, A

ALIGNMENTS

RESULT 1

US-10-355-577-24855/c
; Sequence 24855, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mitmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 24855
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-24855

Query Match 89.3%; Score 13.4; DB 14; Length 25;
Best Local Similarity 73.3%; Pred. No. 1.9e+03;

Matches 11; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGAU 15

Db 25 AGATAAGTCGAGAT 11

RESULT 2

US-60-427-836-243678/c
; Sequence 243678, Application US/60427836
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Rat
; FILE REFERENCE: 3527
; CURRENT APPLICATION NUMBER: US/60/427,836
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 699466

; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 243678
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-60-427-836-243678

Query Match 89.3%; Score 13.4; DB 18; Length 25;
Best Local Similarity 73.3%; Pred. No. 1.9e+03;
Matches 11; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAG 15
DB 21 AGATAACGTGAAGAT 7
|||||1|1|1|1|1|

RESULT 3
US-10-355-577-179244/c
; Sequence 179244, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 179244
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-179244

Query Match 86.7%; Score 13; DB 14; Length 25;
Best Local Similarity 84.6%; Pred. No. 3.1e+03;
Matches 11; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 GAUACGUGAAGA 14
DB 22 GATAACGTGAAGA 10
|||||1|1|1|1|1|

RESULT 4
US-60-427-836-118114
; Sequence 118114, Application US/60427836
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Rat
; FILE REFERENCE: 3527
; CURRENT APPLICATION NUMBER: US/60/427,836
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 699466
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 118114
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-60-427-836-118114

Query Match 86.7%; Score 13; DB 18; Length 25;
Best Local Similarity 84.6%; Pred. No. 3.1e+03;
Matches 11; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAG 13
DB 1 AGATAACGTGAAG 13
|||||1|1|1|1|1|

RESULT 5
US-10-310-188-75880/c
; Sequence 75880, Application US/10310188
; GENERAL INFORMATION:
; APPLICANT: RosettaGenomics

; TITLE OF INVENTION: BIOINFORMATICAALLY DETECTABLE GROUP OF NOVEL VIRAL REGULATORY
; TITLE OF INVENTION: USES THEREOF
; FILE REFERENCE: 47487
; CURRENT APPLICATION NUMBER: US/10/310,188
; CURRENT FILING DATE: 2002-12-19
; NUMBER OF SEQ ID NOS: 86841
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 75880
; LENGTH: 23
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-310-188-75880

Query Match 82.7%; Score 12.4; DB 14; Length 23;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGA 14
DB 23 AAATAACGTGAAGA 10
|1|1|1|1|1|1|1|

RESULT 6
US-09-660-222-72811/c
; Sequence 72811, Application US/09660222
; GENERAL INFORMATION:
; APPLICANT: Mittmann et al.
; TITLE OF INVENTION: Methods of Genetic Analysis of Human
; FILE REFERENCE: 3102.1
; CURRENT APPLICATION NUMBER: US/09/660,222
; CURRENT FILING DATE: 2000-09-12
; PRIOR APPLICATION NUMBER: 60/164,973
; PRIOR FILING DATE: 1999-11-11
; NUMBER OF SEQ ID NOS: 140981
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 72811
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo Sapiens
; PUBLICATION INFORMATION:
; DATABASE ACCESSION NUMBER: GenBank S80343
US-09-660-222-72811

Query Match 82.7%; Score 12.4; DB 10; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGA 14
DB 16 AGATAACGTGCAGA 3
|||||1|1|1|1|1|

RESULT 7
US-09-660-222-72812/c
; Sequence 72812, Application US/09660222
; GENERAL INFORMATION:
; APPLICANT: Mittmann et al.
; APPLICANT: Affymetrix, Inc.
; TITLE OF INVENTION: Methods of Genetic Analysis of Human
; FILE REFERENCE: 3102.1
; CURRENT APPLICATION NUMBER: US/09/660,222
; CURRENT FILING DATE: 2000-09-12
; PRIOR APPLICATION NUMBER: 60/164,973
; PRIOR FILING DATE: 1999-11-11
; NUMBER OF SEQ ID NOS: 140981
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 72812
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo Sapiens
; PUBLICATION INFORMATION:
; DATABASE ACCESSION NUMBER: GenBank S80343

US-09-660-222-72812

Query Match 82.7%; Score 12.4; DB 10; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels

QY 1 AGAUAAACGUGAAGA 14
|||:||||:| |||
Db 22 AGATAACGTGCAGA 9

RESULT 8

```

US-09-953-570-48322/c
; Sequence 48322, Application US/09953570
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic An
; FILE REFERENCE: 310.1
; CURRENT APPLICATION NUMBER: US/09/953,570
; CURRENT FILING DATE: 2001-09-13
; PRIOR APPLICATION NUMBER: 60/232,638
; PRIOR FILING DATE: 2000-09-14
; NUMBER OF SEQ IDS: 138410
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 48322
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Saccharomyces Cerevisiae
US-09-953-570-48322

```

Query Match	82.7%;	Score 12.4;	DB 11;	Length 25;
Best Local Similarity	78.6%;	Pred. No. 6.7e+03;		
Matches 11;	Conservative	2;	Mismatches 1;	Indels

QY 1 AGAUAACGUGAAGA 14
|||:|||||
Db 14 AGATAACGTGTAGA 1

RESULT 9

```

US-09-953-570-48329/c
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; Sequence 48329, Application US/09953570
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic An
; FILE REFERENCE: 3110.1
; CURRENT APPLICATION NUMBER: US/09/953,570
; CURRENT FILING DATE: 2001-09-13
; PRIOR APPLICATION NUMBER: 60/232,638
; PRIOR FILING DATE: 2000-09-14
; NUMBER OF SEQ ID NOS: 138410
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 48329
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Saccharomyces Cerevisiae
US-09-953-570-48329

```

Query Match	82.7%;	Score 12.4;	DB 11;	Length 25;
Best Local Similarity	78.6%;			
Matches 11;	Conservative	2;	Mismatches 1;	Indels 0

QY 1 AGAUAACGUGAAGA 14
 |||:||||:| |||
 Db 20 AGATAACGTGTAGA 7

RESULT 10

US-09-954-445A-10200/c
; Sequence 10200, application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic

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: FILE REFERENCE: 3116.1
: CURRENT APPLICATION NUMBER: US/09/954,445A
: CURRENT FILING DATE: 2000-09-17
: PRIORITY APPLICATION NUMBER: 60/233,620
: PRIOR FILING DATE: 2000-09-18
: NUMBER OF SEQ ID NOS: 131820
: SOFTWARE: Microarray Probe Sequence Listing
: SEQ ID NO 10200
: LENGTH: 25
: TYPE: DNA
: ORGANISM: Arabidopsis thaliana
US-09-954-445A-10200

```

Query Match	82.78;	Score 12.4;	DB 11;	Length 25;
Best Local Similarity	78.63;	Pred. No. 6.7e+03;		
Matches 11; Conservative	2;	Mismatches 1;	Indels 0;	Gaps 0;
Qy	1	AGAAACGUGAAGA	14	
Db	17	AGATAAAGTGAAGA	4	

RESULT 11

```

US-09-954-445A-16612
; Sequence 16612, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954,445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ. ID. NO 16612
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-16612

```

Query Match	82.7%	Score 12.4;	DB 11;	Length 25;
Best Local Similarity	71.4%;	Pred. No. 6.7e+03;		
Matches	10;	Conservative	3;	Mismatches 1;
				Indels 0;
				Gaps 0;
QY	2	GAUACGUGAAGAU	15	
Db	6	GATAACGTGAATAT	19	

RESULT 12

```

US-09-954-445A-16620
; Sequence 16620, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954,445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator v 1.1
; SEQ ID NO 16620
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-16620

```

Query Match	82.7%	Score 12.4	DB 11	Length 25
Best Local Similarity	71.4%	Pred. No. 6.7e+03		
Matches 10	Conservative	3	Mismatches 1	Indels 0
				Gaps 0

; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana

QY 2 GAUAACGUGAAGAU 15
||:||||:|||||:
Db 1 GATAACGTGAATAT 14

RESULT 13

US-09-954-445A-50048
; Sequence 50048, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954.445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 50048
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-50048

Query Match 82.7%; Score 12.4; DB 11; Length 25;
Best Local Similarity 71.4%; Pred. No. 6.7e+03;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGAU 15
||:||||:|||||:
Db 6 GATTACGTGAATAT 19

RESULT 14

US-10-355-577-40591
; Sequence 40591, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355.577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 40591
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-40591

Query Match 82.7%; Score 12.4; DB 14; Length 25;
Best Local Similarity 85.7%; Pred. No. 6.7e+03;
Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
||| ||||:|||||
Db 2 AGAGAACGTGAAGA 15

RESULT 15

US-10-355-577-366476/c
; Sequence 366476, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355.577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 366476
; LENGTH: 25
; TYPE: DNA

; ORGANISM: Homo sapien
US-10-355-577-366476

Query Match 82.7%; Score 12.4; DB 14; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
|||:||||:|||||
Db 18 AGATAACGTGCAGA 5

Search completed: July 6, 2003, 16:49:13
Job time : 522.455 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:26:16 ; Search time 493.182 seconds
(without alignments)
885.154 Million cell updates/sec

Title: US-09-780-929-97
Perfect score: 15
Sequence: 1 agauacgugaagau 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 897812

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl.*

1: gb_ba.*
2: gb_hgt.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*
15: em_ba.*
16: em_fun.*
17: em_hum.*
18: em_in.*
19: em_mu.*
20: em_om.*
21: em_or.*
22: em_ov.*
23: em_pat.*
24: em_ph.*
25: em_pl.*
26: em_ro.*
27: em_sts.*
28: em_un.*
29: em_vi.*
30: em_htg_hum.*
31: em_htg_inv.*
32: em_htg_other.*
33: em_htg_mus.*
34: em_htg_pln.*
35: em_htg_rod.*
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41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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2	15	100.0	27	6	AX214239	Sequence
3	15	100.0	28	6	AX214237	Sequence
4	15	100.0	28	6	AX214238	Sequence
5	15	100.0	28	6	AX214240	Sequence
6	15	100.0	28	6	AX214241	Sequence
7	15	100.0	28	6	AX214242	Sequence
8	15	100.0	28	6	AX214243	Sequence
9	15	100.0	28	6	AX214244	Sequence
10	15	100.0	28	6	AX214245	Sequence
11	15	100.0	28	6	AX214246	Sequence
12	15	100.0	28	6	AX214247	Sequence
13	15	100.0	28	6	AX214248	Sequence
14	15	100.0	28	6	AX214249	Sequence
15	15	100.0	28	6	AX214250	Sequence
16	15	100.0	28	6	AX214251	Sequence
17	15	100.0	28	6	AX214252	Sequence
18	15	100.0	28	6	AX214253	Sequence
19	15	100.0	28	6	AX214254	Sequence
20	15	100.0	28	6	AX214255	Sequence
21	15	100.0	28	6	AX214256	Sequence
22	15	100.0	28	6	AX214257	Sequence
23	15	100.0	28	6	AX214258	Sequence
24	15	100.0	28	6	AX214259	Sequence
25	15	100.0	28	6	AX214260	Sequence
26	15	100.0	28	6	AX214261	Sequence
27	15	100.0	28	6	AX214262	Sequence
28	15	100.0	28	6	AX214263	Sequence
29	15	100.0	28	6	AX214264	Sequence
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ALIGNMENTS

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LOCUS AX214295
DEFINITION Sequence 108 from Patent WO0159102.
ACCESSION AX214295
VERSION AX214295.1 GI:15524372
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 15)
AUTHORS Breaker, R. and Emilsson, G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 108 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)

linear PAT 06-SEP-2001

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  DEFINITION Sequence 52 from Patent WO0159102.
  ACCESSION AX214239
  VERSION AX214239.1 GI:15524316
  KEYWORDS
    SOURCE
      synthetic construct.
      synthetic construct
      artificial sequences.
    ORGANISM
      Breaker,R. and Emilsson,G.
  REFERENCE 1 (bases 1 to 27)
  AUTHORS Nucleozymes with endonuclease activity
  TITLE Patent: WO 0159102-A 52 16-AUG-2001;
  JOURNAL RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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  LOCUS AX214237 28 bp mRNA linear PAT 06-SEP-2001
  DEFINITION Sequence 50 from Patent WO0159102.
  ACCESSION AX214237
  VERSION AX214237.1 GI:15524314
  KEYWORDS
    SOURCE
      synthetic construct.
      synthetic construct
      artificial sequences.
    ORGANISM
      Breaker,R. and Emilsson,G.
  REFERENCE 1 (bases 1 to 28)
  AUTHORS Nucleozymes with endonuclease activity
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  DEFINITION Sequence 51 from Patent WO0159102.
  ACCESSION AX214238
  VERSION AX214238.1 GI:15524315
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    SOURCE
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      synthetic construct
      artificial sequences.
    ORGANISM
      Breaker,R. and Emilsson,G.
  REFERENCE 1 (bases 1 to 28)
  AUTHORS Nucleozymes with endonuclease activity
  TITLE Patent: WO 0159102-A 51 16-AUG-2001;
  JOURNAL RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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  DEFINITION Sequence 53 from Patent WO0159102.
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  VERSION AX214240.1 GI:15524317
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      synthetic construct
      artificial sequences.
    ORGANISM
      Breaker,R. and Emilsson,G.
  REFERENCE 1 (bases 1 to 28)
  AUTHORS Nucleozymes with endonuclease activity
  TITLE Patent: WO 0159102-A 53 16-AUG-2001;
  JOURNAL RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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Db 7 AGATAACGTGAAGAT 21
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LOCUS AX214241 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 54 from Patent WO0159102.
ACCESSION AX214241
VERSION AX214241.1 GI:15524318
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 54 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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Db 7 AGATAACGTGAAGAT 21
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AX214242
LOCUS AX214242 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 55 from Patent WO0159102.
ACCESSION AX214242
VERSION AX214242.1 GI:15524319
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 55 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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Db 7 AGATAACGTGAAGAT 21
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LOCUS AX214243 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 56 from Patent WO0159102.
ACCESSION AX214243
VERSION AX214243.1 GI:15524320
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 56 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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AX214244
LOCUS AX214244 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 57 from Patent WO0159102.
ACCESSION AX214244
VERSION AX214244.1 GI:15524321
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 57 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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AX214245 28 bp mRNA linear PAT 06-SEP-2001
LOCUS
DEFINITION Sequence 58 from Patent WO0159102.
ACCESSION AX214245
VERSION AX214245.1 GI:15524322
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 58 16-AUG-2001.
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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RESULT 11
AX214246 28 bp mRNA linear PAT 06-SEP-2001
LOCUS
DEFINITION Sequence 59 from Patent WO0159102.
ACCESSION AX214246
VERSION AX214246.1 GI:15524323
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequences..

REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 59 16-AUG-2001.
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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DEFINITION Sequence 60 from Patent WO0159102.
ACCESSION AX214247
VERSION AX214247.1 GI:15524324
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 60 16-AUG-2001.
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
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RESULT 13
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LOCUS
DEFINITION Sequence 61 from Patent WO0159102.
ACCESSION AX214248
VERSION AX214248 GI:15524325
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SOURCE      synthetic construct.
ORGANISM     synthetic construct
REFERENCE    1 (bases 1 to 28)
AUTHORS      Breaker,R. and Emilsson,G.
TITLE        Nucleozymes with endonuclease activity
JOURNAL      Patent: WO 0159102-A 61 16-AUG-2001;
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QY 1 AGAUACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

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LOCUS        AX214249          28 bp      mRNA
DEFINITION   Sequence 62 from Patent WO0159102.
ACCESSION    AX214249
VERSION      AX214249.1 GI:15524326
KEYWORDS
SOURCE       synthetic construct.
ORGANISM     synthetic construct
REFERENCE    1 (bases 1 to 28)
AUTHORS      Breaker,R. and Emilsson,G.
TITLE        Nucleozymes with endonuclease activity
JOURNAL      Patent: WO 0159102-A 62 16-AUG-2001;
RIBOZYME     PHARMACEUTICALS, INC. (US) ; Yale University (US)
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SOURCE      synthetic construct.
ORGANISM     synthetic construct
REFERENCE    1 (bases 1 to 28)
AUTHORS      Breaker,R. and Emilsson,G.
TITLE        Nucleozymes with endonuclease activity
JOURNAL      Patent: WO 0159102-A 61 16-AUG-2001;
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RESULT 15
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ORGANISM     synthetic construct
REFERENCE    1 (bases 1 to 28)
AUTHORS      Breaker,R. and Emilsson,G.
TITLE        Nucleozymes with endonuclease activity
JOURNAL      Patent: WO 0159102-A 63 16-AUG-2001;
RIBOZYME     PHARMACEUTICALS, INC. (US) ; Yale University (US)
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Db 7 AGATAACGTGAAGAT 21

Search completed: July 6, 2003, 14:51:11
Job time : 493.182 secs
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:25:15 ; Search time 160.909 Seconds
(without alignments)
209.932 Million cell updates/sec

Title: US-09-780-929-97
Perfect score: 15
Sequence: 1 agaaacgugaagau 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues
Total number of hits satisfying chosen parameters: 2274872

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
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- 13: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.*
- 14: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.*
- 15: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.*
- 16: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.*
- 17: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.*
- 18: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
- 19: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.*
- 20: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.*
- 21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
- 22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
- 23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
- 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	100.0	15	22	AA12347
2	15	100.0	27	22	AA12295
3	15	100.0	27	22	AA12296
4	15	100.0	27	22	AA12297
5	15	100.0	27	22	AA12298
6	15	100.0	27	22	AA12299
7	15	100.0	27	22	AA12300
8	15	100.0	27	22	AA12301
9	15	100.0	27	22	AA12302

10	15	100.0	27	22	AA12303	DNA encoding class
11	15	100.0	27	22	AA12304	DNA encoding class
12	15	100.0	27	22	AA12305	DNA encoding class
13	15	100.0	27	22	AA12306	DNA encoding class
14	15	100.0	27	22	AA12307	DNA encoding class
15	15	100.0	27	22	AA12308	DNA encoding class
16	15	100.0	27	22	AA12309	DNA encoding class
17	15	100.0	27	22	AA12310	DNA encoding class
18	15	100.0	27	22	AA12311	DNA encoding class
19	15	100.0	27	22	AA12312	DNA encoding class
20	15	100.0	27	22	AA12313	DNA encoding class
21	15	100.0	27	22	AA12314	DNA encoding class
22	15	100.0	27	22	AA12315	DNA encoding class
23	15	100.0	27	22	AA12316	DNA encoding class
24	15	100.0	27	22	AA12317	DNA encoding class
25	15	100.0	27	22	AA12318	DNA encoding class
26	15	100.0	27	22	AA12319	DNA encoding class
27	15	100.0	27	22	AA12320	DNA encoding class
28	15	100.0	27	22	AA12321	DNA encoding class
29	15	100.0	27	22	AA12322	DNA encoding class
30	15	100.0	27	22	AA12323	DNA encoding class
31	15	100.0	27	22	AA12324	DNA encoding class
32	15	100.0	27	22	AA12325	DNA encoding class
33	15	100.0	27	22	AA12326	DNA encoding class
34	15	100.0	27	22	AA12327	DNA encoding class
35	15	100.0	27	22	AA12328	DNA encoding class
36	15	100.0	27	22	AA12329	DNA encoding class
37	15	100.0	27	22	AA12330	DNA encoding class
38	15	100.0	27	22	AA12331	DNA encoding class
39	15	100.0	27	22	AA12332	DNA encoding class
40	15	100.0	27	22	AA12333	DNA encoding class
41	15	100.0	27	22	AA12334	DNA encoding class
42	15	100.0	27	22	AA12335	DNA encoding class
43	15	100.0	27	22	AA12336	DNA encoding class
44	15	100.0	27	22	AA12337	DNA encoding class
45	15	100.0	27	22	AA12404	DNA encoding class

ALIGNMENTS

RESULT 1

AA12347
ID AA12347 standard; DNA; 15 BP.

XX AA12347;

XX 21-NOV-2001 (first entry)

XX DNA encoding deoxyribozyme #7.

XX Deoxyribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ribozyme; ss.

XX Synthetic.

XX WO200159102-A2.

XX 16-AUG-2001.

XX 08-FEB-2001; 2001WO-US04223.

XX 08-FEB-2000; 2000US-0181360.

XX 31-MAR-2000; 2000US-0193646.

XX (RIBO-) RIBOZYME PHARM INC.

XX (UYFA) UNIV YALE.

XX Breaker R, Beigelman L, Emilsson G;

XX WPI; 2001-536526/59.

XX New nucleic acids with endonuclease activity, such as ribozymes and

PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell
XX
PS Claim 49; Page 77; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of deoxyribozyme #7 used in the method of the invention.
XX
SQ Sequence 15 BP; 7 A; 1 C; 4 G; 3 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 15;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
|||||
Db 1 AGAUAACGUGAAGAU 15

RESULT 2

AAS12295
ID AAS12295 standard; DNA; 27 BP.

AC AAS12295;

XX 21-NOV-2001 (first entry)

XX DNA encoding class V ribozyme #7.

XX Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.

XX Synthetic.

XX Key Location/Qualifiers

FT modified_base 1..4
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 23..27
FT /*tag= b
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 27
FT /*tag= c
FT /mod_base= c
FT /note= "3',3'-inverted deoxybasic moiety"

XX WO200159102-A2.

XX 16-AUG-2001.

XX 08-FEB-2001; 2001WO-US04223.

XX 08-FEB-2000; 2000US-0181360.

XX 31-MAR-2000; 2000US-0193646.

XX (RIBO-) RIBOZYME PHARM INC.
XX (UYIA) UNIV YALE.

XX Breaker R, Beigelman L, Emilsson G;

DR WPI; 2001-536526/59.
XX
PT New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell
XX
PS Example 1; Page 71; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #7 used in the method of the invention.
XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
|||||
Db 7 AGAUAACGUGAAGAU 21

RESULT 3

AAS12296

ID AAS12296 standard; DNA; 27 BP.

AC AAS12296;

XX 21-NOV-2001 (first entry)

XX DNA encoding class V ribozyme #8.

XX Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.

XX Synthetic.

XX Key Location/Qualifiers

FT modified_base 1..6
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 20..27
FT /*tag= b
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 27
FT /*tag= c
FT /mod_base= c
FT /note= "3',3'-inverted deoxybasic moiety"

XX WO200159102-A2.

XX 16-AUG-2001.

XX 08-FEB-2001; 2001WO-US04223.

XX 08-FEB-2000; 2000US-0181360.

XX 31-MAR-2000; 2000US-0193646.

XX (RIBO-) RIBOZYME PHARM INC.
XX (UYIA) UNIV YALE.

```
XX PI Breaker R, Beigelman L, Emilsson G;
XX PA WPI; 2001-536526/59.
XX DR
XX XX
XX PT New nucleic acids with endonuclease activity, such as ribozymes and
XX FT nucleozymes, for modulating gene expression in a plant, mammalian,
XX FT bacterial or fungal cell
XX PS
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #8 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 4
AAS12297
ID AAS12297 standard; DNA; 27 BP.
XX AC AAS12297;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #9.
XX KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /*tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 21..27
XX FT /*tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /*tag= c
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxyabasic moiety"
XX PN WO200159102-A2.
XX PD 16-AUG-2001.
XX PF 08-FEB-2001; 2001WO-US04223.
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
```

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XX XX (RIBO-) RIBOZYME PHARM INC.
XX PA (UYVA ) UNIV YALE.
XX XX
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX
XX DR WPI; 2001-536526/59.
XX XX
XX PT New nucleic acids with endonuclease activity, such as ribozymes and
XX FT nucleozymes, for modulating gene expression in a plant, mammalian,
XX FT bacterial or fungal cell
XX PS
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #9 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 5
AAS12298
ID AAS12298 standard; DNA; 27 BP.
XX AC AAS12298;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #10.
XX KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /*tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 12
XX FT /*tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 21..27
XX FT /*tag= c
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /*tag= d
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxyabasic moiety"
XX PN WO200159102-A2.
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XX PD 16-AUG-2001.
XX XX
XX PF 08-FEB-2001; 2001WO-US04223.
XX XX
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX XX
XX PA (RIBO-) RIBOZYME PHARM INC.
XX PA (UYIA ) UNIV YALE.
XX XX
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX
XX DR WPI; 2001-536526/59.
XX XX
XX PT New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell -
XX XX
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #10 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGUAACGUGAAGAU 15
Db 7 AGUAACGUGAAGAU 21

RESULT 6
AAS12299
ID AAS12299 standard; DNA; 27 BP.
XX AC AAS12299;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #11.
XX KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..7
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 21..27
XX FT /tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= c
XX FT /mod_base= c

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FT XX /note= "3',3'-inverted deoxyabasic moiety"
XX PN WO200159102-A2.
XX XX
XX PD 16-AUG-2001.
XX XX
XX PF 08-FEB-2001; 2001WO-US04223.
XX XX
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX XX
XX XX (RIBO-) RIBOZYME PHARM INC.
XX PA (UYIA ) UNIV YALE.
XX XX
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX
XX DR WPI; 2001-536526/59.
XX XX
XX PT New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell -
XX XX
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #11 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGUAACGUGAAGAU 15
Db 7 AGUAACGUGAAGAU 21

RESULT 7
AAS12300
ID AAS12300 standard; DNA; 27 BP.
XX AC AAS12300;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #12.
XX KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 21..27
XX FT /tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"

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```
FT modified_base 27 /*tag= c
FT /*mod_base= c
FT /*note= "3',3'-inverted deoxybasic moiety"
XX
PN WO200159102-A2.
XX
PD 16-AUG-2001.
XX
PF 08-FEB-2001; 2001WO-US04223.
XX
PR 08-FEB-2000; 2000US-0181360.
PR 31-MAR-2000; 2000US-0193646.
XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (UYVA ) UNIV YALE.
XX
PI Breaker R, Beigelman L, Emilsson G;
XX
DR WPI; 2001-536526/59.
XX
PS New nucleic acids with endonuclease activity, such as ribozymes and
XX bacterial or fungal cell
XX
XX Example 1; Page 71; 96pp; English.
XX
XX The invention relates to nucleic acid molecules with endonuclease
XX activity, which are particularly useful for cleavage of RNA or DNA.
XX The nucleic acids are used in a pharmaceutical composition and are used
XX to modulate expression of a gene in a plant, mammalian, bacterial or
XX fungal cell. They are used to cleave a separate nucleic acid, preferably
XX RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX proliferation, and can be used to treat a disease or condition. More
XX than one nucleic acid can be independently targeted to the same or
XX different sites in a cell. The nucleic acids may be used to study DNA.
XX The modifications to the nucleic acids optimises their catalytic activity
XX and can maintain or enhance their activity. They exhibit a high degree
XX of specificity for RNA. The present sequence represents the coding
XX sequence of class V ribozyme #12 used in the method of the invention.
XX
XX Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX
XX Query Match 100.0%; Score 15; DB 22; Length 27;
XX Best Local Similarity 100.0%; Pred. No. 74;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 AGAUAACGUGAAGAU 15
XX |||||
XX Db 7 AGAUAACGUGAAGAU 21
XX
XX RESULT 8
XX AAS12301
XX ID AAS12301 standard; DNA; 27 BP.
XX
XX AC AAS12301;
XX
XX DT 21-NOV-2001 (first entry)
XX
XX DE DNA encoding class V ribozyme #13.
XX
XX KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX
XX OS Synthetic.
XX
XX FH Key Location/Qualifiers
XX modified_base 1..6
XX /*tag= a
XX /*mod_base= OTHER
XX /*note= "OTHER = 2'-O-methyl nucleotides"
XX
XX modified_base 18
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```
FT /*tag= b
FT /*mod_base= a
FT /*note= "OTHER = 2'-O-methyl nucleotide"
XX
XX modified_base 21..27
FT /*tag= c
FT /*mod_base= OTHER
FT /*note= "OTHER = 2'-O-methyl nucleotides"
XX
XX modified_base 27
FT /*tag= d
FT /*mod_base= c
FT /*note= "3',3'-inverted deoxybasic moiety"
XX
XX PN WO200159102-A2.
XX
XX PD 16-AUG-2001.
XX
XX PF 08-FEB-2001; 2001WO-US04223.
XX
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX (UYVA ) UNIV YALE.
XX
XX PI Breaker R, Beigelman L, Emilsson G;
XX
XX DR WPI; 2001-536526/59.
XX
XX PS New nucleic acids with endonuclease activity, such as ribozymes and
XX nucleozymes, for modulating gene expression in a plant, mammalian,
XX bacterial or fungal cell
XX
XX Example 1; Page 71; 96pp; English.
XX
XX The invention relates to nucleic acid molecules with endonuclease
XX activity, which are particularly useful for cleavage of RNA or DNA.
XX The nucleic acids are used in a pharmaceutical composition and are used
XX to modulate expression of a gene in a plant, mammalian, bacterial or
XX fungal cell. They are used to cleave a separate nucleic acid, preferably
XX RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX proliferation, and can be used to treat a disease or condition. More
XX than one nucleic acid can be independently targeted to the same or
XX different sites in a cell. The nucleic acids may be used to study DNA.
XX The modifications to the nucleic acids optimises their catalytic activity
XX and can maintain or enhance their activity. They exhibit a high degree
XX of specificity for RNA. The present sequence represents the coding
XX sequence of class V ribozyme #13 used in the method of the invention.
XX
XX Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX
XX Query Match 100.0%; Score 15; DB 22; Length 27;
XX Best Local Similarity 100.0%; Pred. No. 74;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 AGAUAACGUGAAGAU 15
XX |||||
XX Db 7 AGAUAACGUGAAGAU 21
XX
XX RESULT 9
XX AAS12302
XX ID AAS12302 standard; DNA; 27 BP.
XX
XX AC AAS12302;
XX
XX DT 21-NOV-2001 (first entry)
XX
XX DE DNA encoding class V ribozyme #14.
XX
XX KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX
XX OS Synthetic.
```

```
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 17
XX FT /tag= b
XX FT /mod_base= a
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 21..27
XX FT /tag= c
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= d
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxybasic moiety"
XX PN WO200159102-A2.
XX PD 16-AUG-2001.
XX PF 08-FEB-2001; 2001WO-US04223.
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX PA (RIBO-) RIBOZYME PHARM INC.
XX PA (UYVA ) UNIV YALE.
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX WPI; 2001-536526/59.
XX XX New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #14 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX Query Match 100.0%; Score 15; DB 22; Length 27;
XX Best Local Similarity 100.0%; Pred. No. 74;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
|||||
RESULT 10
AAS12303
ID AAS12303 standard; DNA; 27 BP.
XX AC AAS12303;
XX XX 21-NOV-2001 (first entry)
DT
```

```
XX DE DNA encoding class V ribozyme #15.
XX KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 11
XX FT /tag= b
XX FT /mod_base= a
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 16
XX FT /tag= c
XX FT /mod_base= g
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 21..27
XX FT /tag= d
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= e
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxybasic moiety"
XX PN WO200159102-A2.
XX PD 16-AUG-2001.
XX PF 08-FEB-2001; 2001WO-US04223.
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX PA (RIBO-) RIBOZYME PHARM INC.
XX PA (UYVA ) UNIV YALE.
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX WPI; 2001-536526/59.
XX XX New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #15 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX Query Match 100.0%; Score 15; DB 22; Length 27;
XX Best Local Similarity 100.0%; Pred. No. 74;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
```


CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #17 used in the method of the invention.

XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;

Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGUAUACGUGAAGAU 15

Db 7 AGUAUACGUGAAGAU 21

RESULT 13

AAS12306

ID AAS12306 standard; DNA; 27 BP.

XX

AC AAS12306;

XX

DT 21-NOV-2001 (first entry)

XX

DE DNA encoding class V ribozyme #18.

XX Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.

XX Synthetic.

OS

XX

FH Key Location/Qualifiers

FT modified_base 1..6

FT /tag= a

FT /mod_base= OTHER

FT /note= "OTHER = 2'-O-methyl nucleotides"

FT 14

FT /tag= b

FT /mod_base= g

FT /note= "OTHER = 2'-O-methyl nucleotide"

FT 21..27

FT /tag= c

FT /mod_base= OTHER

FT /note= "OTHER = 2'-O-methyl nucleotides"

FT 27

FT /tag= d

FT /mod_base= c

FT /note= "3',3'-inverted deoxyabasic moiety"

FT

XX WO200159102-A2.

XX

XX 16-AUG-2001.

XX

XX 08-FEB-2001; 2001WO-US04223.

XX

XX 08-FEB-2000; 2000US-0181360.

XX

XX 31-MAR-2000; 2000US-0193646.

XX

XX (RIBO-) RIBOZYME PHARM INC.

XX

XX (UYVA) UNIV YALE.

XX

XX Breaker R, Beigelman L, Emilsson G;

XX

XX WPI; 2001-536526/59.

XX

XX New nucleic acids with endonuclease activity, such as ribozymes and

XX nucleozymes, for modulating gene expression in a plant, mammalian,

XX bacterial or fungal cell

XX

XX Example 1; Page 71; 96pp; English.

XX

XX The invention relates to nucleic acid molecules with endonuclease

XX activity, which are particularly useful for cleavage of RNA or DNA.

XX The nucleic acids are used in a pharmaceutical composition and are used

XX to modulate expression of a gene in a plant, mammalian, bacterial or

CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #18 used in the method of the invention.

XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;

Best Local Similarity 100.0%; Pred. No. 74;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGUAUACGUGAAGAU 15

Db 7 AGUAUACGUGAAGAU 21

RESULT 14

AAS12307

ID AAS12307 standard; DNA; 27 BP.

XX

AC AAS12307;

XX

DT 21-NOV-2001 (first entry)

XX

DE DNA encoding class V ribozyme #19.

XX

KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;

KW gene therapy; plant; fungus; bacteria; mammal; ss.

XX

OS Synthetic.

XX

FH Key Location/Qualifiers

FT modified_base 1..6

FT /tag= a

FT /mod_base= OTHER

FT /note= "OTHER = 2'-O-methyl nucleotides"

FT 13

FT /tag= b

FT /mod_base= c

FT /note= "OTHER = 2'-O-methyl nucleotide"

FT 21..27

FT /tag= c

FT /mod_base= OTHER

FT /note= "OTHER = 2'-O-methyl nucleotides"

FT 27

FT /tag= d

FT /mod_base= c

FT /note= "3',3'-inverted deoxyabasic moiety"

FT

XX WO200159102-A2.

XX

XX 16-AUG-2001.

XX

XX 08-FEB-2001; 2001WO-US04223.

XX

XX 08-FEB-2000; 2000US-0181360.

XX

XX 31-MAR-2000; 2000US-0193646.

XX

XX (RIBO-) RIBOZYME PHARM INC.

XX

XX (UYVA) UNIV YALE.

XX

XX Breaker R, Beigelman L, Emilsson G;

XX

XX WPI; 2001-536526/59.

XX

XX New nucleic acids with endonuclease activity, such as ribozymes and

XX nucleozymes, for modulating gene expression in a plant, mammalian,

XX bacterial or fungal cell

XX

XX PS Example 1; Page 71; 96pp; English.

XX CC The invention relates to nucleic acid molecules with endonuclease

CC activity, which are particularly useful for cleavage of RNA or DNA.

CC The nucleic acids are used in a pharmaceutical composition and are used

CC to modulate expression of a gene in a plant, mammalian, bacterial or

CC fungal cell. They are used to cleave a separate nucleic acid, preferably

CC RNA. The nucleic acids are used to inhibit gene expression and/or cell

CC proliferation, and can be used to treat a disease or condition. More

CC than one nucleic acid can be independently targeted to the same or

CC different sites in a cell. The nucleic acids may be used to study DNA.

CC The modifications to the nucleic acids optimises their catalytic activity

CC and can maintain or enhance their activity. They exhibit a high degree

CC of specificity for RNA. The present sequence represents the coding

CC sequence of class V ribozyme #19 used in the method of the invention.

XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;

Best Local Similarity 100.0%; Pred. No. 74;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15

Db 7 AGAUACGUGAAGAU 21

RESULT 15

AAS12308

ID AAS12308 standard; DNA; 27 BP.

XX AC AAS12308;

XX DT 21-NOV-2001 (first entry)

XX DE DNA encoding class V ribozyme #20.

XX KW Ribozyme; cytosstatic; endonuclease; RNA cleavage; DNA cleavage;

XX KW gene therapy; plant; fungus; bacteria; mammal; ss.

XX OS Synthetic.

XX FH Key Location/Qualifiers

FT modified_base 1..6

FT /*tag= a

FT /mod_base= OTHER

FT /note= "OTHER = 2'-O-methyl nucleotides"

FT modified_base 21..27

FT /*tag= b

FT /mod_base= OTHER

FT /note= "OTHER = 2'-O-methyl nucleotides"

FT modified_base 27

FT /*tag= c

FT /mod_base= c

FT /note= "3',3'-inverted deoxybasic moiety"

XX PN WO200159102-A2.

XX PD 16-AUG-2001.

XX PF 08-FEB-2001; 2001WO-US04223.

XX PR 08-FEB-2000; 2000US-0181360.

XX PR 31-MAR-2000; 2000US-0193646.

XX PA (RIBO-) RIBOZYME PHARM INC.

XX PA (UYA) UNIV YALE.

XX PI Breaker R, Beigelman L, Emilsson G;

XX DR WPI; 2001-536526/59.

XX

PT New nucleic acids with endonuclease activity, such as ribozymes and

PT nucleozymes, for modulating gene expression in a plant, mammalian,

PT bacterial or fungal cell

XX Example 1; Page 71; 96pp; English.

XX CC The invention relates to nucleic acid molecules with endonuclease

CC activity, which are particularly useful for cleavage of RNA or DNA.

CC The nucleic acids are used in a pharmaceutical composition and are used

CC to modulate expression of a gene in a plant, mammalian, bacterial or

CC fungal cell. They are used to cleave a separate nucleic acid, preferably

CC RNA. The nucleic acids are used to inhibit gene expression and/or cell

CC proliferation, and can be used to treat a disease or condition. More

CC than one nucleic acid can be independently targeted to the same or

CC different sites in a cell. The nucleic acids may be used to study DNA.

CC The modifications to the nucleic acids optimises their catalytic activity

CC and can maintain or enhance their activity. They exhibit a high degree

CC of specificity for RNA. The present sequence represents the coding

CC sequence of class V ribozyme #20 used in the method of the invention.

XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;

Best Local Similarity 100.0%; Pred. No. 74;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15

Db 7 AGAUACGUGAAGAU 21

Search completed: July 6, 2003, 14:32:52

Job time : 160.909 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:26:51 : Search time 1007.73 Seconds
(without alignments)
241.069 Million cell updates/sec

Title: US-09-780-929-97

Perfect score: 15

Sequence: 1 agaaacgugaagau 15

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 146654

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: em_estba:*

2: em_esthum:*

3: em_estin:*

4: em_estmu:*

5: em_estov:*

6: em_estpl:*

7: em_estro:*

8: em_htc:*

9: gb_estl:*

10: gb_est2:*

11: gb_htc:*

12: gb_est3:*

13: gb_est4:*

14: gb_est5:*

15: em_estfun:*

16: em_estom:*

17: gb_gss:*

18: em_gss_hum:*

19: em_gss_inv:*

20: em_gss_pln:*

21: em_gss_rft:*

22: em_gss_fun:*

23: em_gss_mam:*

24: em_gss_mus:*

25: em_gss_other:*

26: em_gss_pro:*

27: em_gss_rod:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
C 1	11.8	78.7	29	17 AZ994980	AZ994980 2M0280G07
2	11.8	78.7	59	14 BQ625912	BQ625912 ph86f02.y
C 3	10.8	72.0	22	17 TA248D01P	AL483187 T. brucei
4	10.8	72.0	28	17 AZ345853	AZ345853 IM0080F14
5	10.8	72.0	37	9 AA947987	AA947987 oq58e02.s
6	10.8	72.0	37	17 AL760544	AL760544 Arabidops

C	7	10.8	72.0	40	9	AI039249	AI039249	ox33a08..s
8	10.8	72.0	46	17	AZ482955	AZ482955	IM0308M13	
9	10.8	72.0	48	17	AZ656367	AZ656367	IM0531D23	
10	10.8	72.0	48	17	AZ772291	AZ772291	IM0583L06	
C	11	10.8	72.0	50	9	AU103231	AU103231	IM0583L06
12	10.8	72.0	51	10	AV833384	AV833384	AV833384	
13	10.8	72.0	52	9	AA823664	AA823664	vr69009..s	
C	14	10.8	72.0	52	12	BF631905	BF631905	NF016H07D
15	10.8	72.0	53	9	AU257573	AU257573	AU257573	
C	16	10.8	72.0	54	13	BM069548	BM069548	ie89d12..x
17	10.8	72.0	54	17	AF149434	AF149434	AF149434	
C	18	10.8	72.0	58	9	AA780323	AA780323	af52b09..s
19	10.8	72.0	59	13	BJ053305	BJ053305	BJ053305	
20	10.4	69.3	42	13	BJ044345	BJ044345	BJ044345	
21	10.4	69.3	44	14	D18694	D18694	MUSGS01756	
22	10.4	69.3	48	14	D18688	D18688	MUSGS01750	
23	10.4	69.3	51	10	AW100987	AW100987	sd64b07..y	
24	10.4	69.3	53	17	AL764237	AL764237	Arabidops	
C	25	10.4	69.3	55	10	AW424064	AW424064	sh60c04..y
26	10.4	69.3	56	12	BG731546	BG731546	dac26g09..	
C	27	10.4	69.3	56	17	AZ784261	AZ784261	2M0026H22
28	10.4	69.3	57	9	AU256712	AU256712	AU256712	
29	10.4	69.3	58	9	AA265174	AA265174	mz49h04..r	
C	30	10.4	69.3	60	13	BI493320	BI493320	df99g12..y
C	31	10.4	69.3	60	17	AL752372	AL752372	Arabidops
32	10.2	68.0	19	17	AZ759898	AZ759898	IM0553A08	
C	33	10.2	68.0	24	17	TA200H11P	TA200H11P	T..brucei
34	10.2	68.0	33	17	AL760055	AL760055	Arabidops	
C	35	10.2	68.0	37	17	AZ760010	AZ760010	IM0553D21
C	36	10.2	68.0	40	9	AI609205	AI609205	tw83a05..x
C	37	10.2	68.0	40	9	AI638565	AI638565	ts50b02..x
38	10.2	68.0	40	14	N54453	N54453	Yv40a04..sl	
39	10.2	68.0	42	17	AZ388234	AZ388234	IM0148E12	
C	40	10.2	68.0	42	17	AZ449920	AZ449920	IM0248I20
C	41	10.2	68.0	44	12	BE738334	BE738334	601572718
42	10.2	68.0	45	17	BH638406	BH638406	IM00802D0	
C	43	10.2	68.0	46	9	AI077563	AI077563	oz33q04..x
C	44	10.2	68.0	46	17	AZ321153	AZ321153	IM0041M11
C	45	10.2	68.0	48	17	BH863621	BH863621	SALK 0942

ALIGNMENTS

RESULT 1
AZ994980/c
LOCUS
DEFINITION 2M0280G07R Mouse 10kb plasmid UUC2M library Mus musculus genomic clone UUC2M0280G07 R, DNA sequence.
ACCESSION AZ994980
VERSION AZ994980.1 GI:13866207
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 29)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A., and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: rdunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00

Plate: 0280 row: G column: 07
 Seq primer: CACACGGAACACCTATGACC
 Class: plasmid ends
 High quality sequence stop: 29.
 Location/Qualifiers

FEATURES

source

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1. .29
  /organism="Mus musculus"
  /strain="C57BL/6J"
  /db_xref="taxon:10090"
  /clone="UUGC2M0280G07"
  /clone_lib="Mouse 10kb plasmid UUGC2M library"
  /sex="Female"
  /lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
  /note="Vector: PWD42nv; Purified genomic DNA from M. Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gil4732114[gb]AF129072.1), a copy-number inducible derivative of plasmid RL. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor-mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
```

```
5 a      5 g      14 t
BASE COUNT
ORIGIN
```

```
Query Match      78.7%; Score 11.8; DB 17; Length 29;
Best Local Similarity 66.7%; Pred. No. 1.2e+04;
Matches 10; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
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Qy 1 AGAUACGUGAAGAU 15

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||||| 1:|||||
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Db 23 AGATAGAGTGAAGAT 9

RESULT 2

BQ625912

LOCUS

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DEFINITION      BQ625912      59 bp      mRNA      linear      EST 01-JUL-2002
                  ph86f02.v1 Ostertagia ostertagi L3 SL1 TOPO v2 Ostertagia ostertagi
                  cDNA 5', mRNA sequence.
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ACCESSION BQ625912

VERSION BQ625912.1

KEYWORDS GI:21653090

SOURCE EST

ORGANISM Ostertagia ostertagi.

```
REFERENCE      Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida;
                  Trichostrongylidae; Haemonchidae; Ostertagiinae; Ostertagia.
```

AUTHORS 1 (bases 1 to 59)

McCartner,J., Clifton,S., Chiapelli,B., Pape,D., Martin,J., Wylie,T.,
 Dante,M., Marra,M., Hillier,L., Kucaba,T., Theising,B., Bowers,Y.,
 Gibbons,M., Ritter,E., Bennett,J., Franklin,C., Tsagarisvili,R.,
 Ronko,I., Kennedy,S., Maguire,L., Beck,C., Underwood,K., Steptoe,
 M., Allen,M., Person,B., Swaller,T., Harvey,N., Schurk,R., Kohn,S.,
 Shin,T., Jackson,Y., Cardenas,M., McCann,R., Waterston,R. and
 Wilson,R.

The Washington Univ. Nematode EST Project, 1999

Unpublished (1999)

Contact: McCarter JP

The Washington Univ. Nematode EST Project, 1999

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. Nematodes were provided by Dr. Louis Gasbarre of the USDA, Beltsville, MD (lgasbarre@nri.barc.usda.gov).
 Putative full length read
 The vector to vector length is 60

Seq primer: SL1 primer.

FEATURES

source

Location/Qualifiers

1. .59

/organism="Ostertagia ostertagi"

/db_xref="taxon:6317"

/clone_lib="Ostertagia ostertagi L3 SL1 TOPO v2"

/dev_stage="third stage exsheathed larvae"

/lab_host="DH10B"

/note="Vector: pCRII-TOPO (Invitrogen); Site_1: EcoRI;

Site_2: EcoRI; The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. Oligo(dT)-SL1 PCR based library. Ostertagia ostertagi L3 cDNA PCR products of size >400 nucleotides containing SL1 on the 5' end and oligo(dT) on the 3' end were non-directionally cloned into pCRII-TOPO(Invitrogen) following the ToPO TA cloning protocol. Nematodes were provided by Dr. Louis Gasbarre of the USDA, Beltsville, MD (lgasbarre@nri.barc.usda.gov). Third stage exsheathed larvae were collected from 14 day fecal-sphagnum moss cultures of Ostertagia eggs. The larvae were recovered by overnight passage on a Baermann apparatus, and then cleaned by passage through a 20 micron nylon mesh. The larvae were then subjected to a treatment with 1.25% chlorox to induce excystation. The larvae were washed with 5 changes of PBS and then pelleted and snap frozen in liquid nitrogen."

BASE COUNT 22 a 16 c 12 g 9 t

ORIGIN

```
Query Match      78.7%; Score 11.8; DB 14; Length 59;
Best Local Similarity 73.3%; Pred. No. 1.7e+04;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
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Qy 1 AGAUACGUGAAGAU 15

```
||||| 1:|||||
```

Db 7 AGAAAAGGTGAAGAT 21

RESULT 3

TA248D01P/c

LOCUS

```
DEFINITION      TA248D01P      22 bp      DNA      linear      GSS 13-DEC-2000
                  T. brucei sheared genomic DNA clone 248d01, forward sequence,
                  genomic survey sequence.
```

ACCESSION AL483187

VERSION AL483187.1

KEYWORDS GI:11848863

SOURCE GSS

ORGANISM Trypanosoma brucei.

Trypanosoma brucei

Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;

Trypanosoma.

REFERENCE 1 (bases 1 to 22)

AUTHORS

Hall,N., Bowman,S., Lennard,N.J., Doggett,J., Atkin,R.,

Chillingworth,C., Ormond,D., Harris,B., El-Sayed,N., Hou,L.,

Melville,S.E., Rajandream,M.A. and Barrell,B.G.

Direct Submission

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
 Project, Sanger Centre, The Wellcome Trust Genome Campus,
 Cambridge CB10 1SA, E-mail: barrellesanger.ac.uk and
 nhlesanger.ac.uk

COMMENT

Constructed at the Institute for Genomic Research (TIGR),
 Rockville, MD. Genomic DNA isolated from a cloned population of
 Trypanosoma brucei (TREU927/4 GUTAT 10.1) was mechanically sheared
 to give a tight size distribution (4 kb). The v + i method used for the library construction is
 described in detail in Smith, H. and Venter, J.C. (Making small
 insert libraries for whole genome shotgun sequencing projects. In
 Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.

Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available
at http://www.sanger.ac.uk/Projects/T_brucei/.

FEATURES

Location/Qualifiers
1..22
/organism="Trypanosoma brucei"
/strain="TRE927"
/db_xref="taxon:5691"
/clone="248d01"

BASE COUNT
ORIGIN

5 a 6 c 2 g 9 t

Query Match 72.0%; Score 10.8; DB 17; Length 22;
Best Local Similarity 64.3%; Pred. No. 3.8e+04;
Matches 9; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY

2 GAUAACGUGAAGAU 15

DB 19 GATACATGAAAT 6

RESULT 4

AZ345853

LOCUS

DEFINITION AZ345853 28 bp DNA linear GSS 29-SEP-2000
clone UUGC1M0080F14 R, DNA sequence.

ACCESSION AZ345853

VERSION AZ345853.1 GI:10425090

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 28)

AUTHORS

Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

JOURNAL Unpublished (2000)

COMMENT

Contact: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0080 row: F column: 14

Seq primer: CACACAGGAACAGCTATGACC

Class: plasmid ends

High quality sequence stop: 28.

FEATURES

Location/Qualifiers
1..28

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC1M0080F14"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(<http://www.jax.org/resources/documents/dnares/>). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (g114732114|g14732114|g14732114), a copy-number
inducible derivative of plasmid RL. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance.

BASE COUNT 16 a 3 c 4 g 5 t

ORIGIN

Query Match 72.0%; Score 10.8; DB 17; Length 28;
Best Local Similarity 71.4%; Pred. No. 4.3e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY

1 AGAUAACGUGAAGA 14

DB 9 AAATAACGTGAAA 22

RESULT 5

AA947987

LOCUS

DEFINITION AA947987 37 bp mRNA linear EST 04-MAY-1998
Oq58e02.s1 NCI-CGAP_Kid6 Homo sapiens cDNA clone IMAGE:1590554 3'
similar to TR:Q33559 Q33559 NH2 TERMINUS UNCERTAIN ;, mRNA

ACCESSION AA947987

VERSION AA947987.1 GI:3109240

KEYWORDS EST..

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 37)

AUTHORS

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

JOURNAL Unpublished (1997)

COMMENT

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: Stratagene, Inc.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone Distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 1.

FEATURES

Location/Qualifiers
1..37

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1590554"

/clone_lib="NCI-CGAP_Kid6"

/sex="mixed"

/tissue_type="kidney tumor"

/lab_host="SOLR (kanamycin resistant)"

/note="Organ: Kidney; Vector: Bluescript SK-; Site:1;

ECORI; Site:2: XhoI; Cloned unidirectionally. Primer:

Oligo dr. Pooled kidney tumors. 5' adaptor sequence: 5'

GAATTCGGCAGAG 3' 3' adaptor sequence: 5'

CTCGAGTTTTTTTTTTTTTTT 3' Average insert size: 1.0 kb.

BASE COUNT 26 a 2 c 6 g 3 t

ORIGIN

Query Match 72.0%; Score 10.8; DB 9; Length 37;
Best Local Similarity 78.6%; Pred. No. 4.9e+04;

```

Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAAUACGUGAAGA 14
Db 17 AGATAATGAGAAGA 30

RESULT 6
AL760544
LOCUS
DEFINITION
Arabidopsis thaliana T-DNA flanking sequence GK-199B11-014765,
genomic survey sequence.
ACCESSION
AL760544
VERSION
AL760544.1 GI:21499415
KEYWORDS
GSS.
SOURCE
thale cress.
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1
REFERENCE
AUTHORS
Strizhov,N., Li,Y., Rosso,M., Viehoever,P., Dekker,K., Saedler,H.
and Weissshaar,B.
TITLE
A pipeline for automated high-throughput generation of ESTs
(flanking sequence tags) from Arabidopsis thaliana T-DNA
transformed lines
JOURNAL
Unpublished
2
REFERENCE
AUTHORS
Rosso,M., Strizhov,N., Li,Y., Reiss,B., Dekker,K. and Weissshaar,B.
TITLE
A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)
for flanking sequence tag based reverse genetics
JOURNAL
Unpublished
3
REFERENCE
AUTHORS
Li,Y., Strizhov,N., Rosso,M. and Weissshaar,B.
TITLE
Direct Submission
JOURNAL
Submitted (17-JUN-2002) Weissshaar B., Max-Planck-Institut fuer
Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
COMMENT
This sequence is recovered from the left border of the T-DNA. It
indicates an insertion close to or within gene Atig77800. The
sequences are generated at the MPI for Plant Breeding Research in
the context of the GABI-Kat project. GABI-Kat is part of the German
Plant Genomics program designated 'GABI'. Information on line
availability can be found at:
http://www.mpiz-koeln.mpg.de/GABI-Kat/.
FEATURES
Location/Qualifiers
1..37
/organism="Arabidopsis thaliana"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-199B11-014765"
/note="PCR was performed on DNA from Arabidopsis thaliana
plants (T1) which were transformed with the T-DNA from
vector PAC161. The lines contain one or more T-DNA
insertions. The DNA fragment(s) resulting from the PCR
were directly sequenced to determine the genomic sequence
flanking the insertion. Sequences displaying significant
similarity to the A. thaliana nuclear genome sequence were
processed for submission. T-DNA derived sequences were
removed"
BASE COUNT 19 a 4 c 6 g 8 t
ORIGIN
Query Match 72.0%; Score 10.8; DB 17; Length 37;
Best Local Similarity 78.6%; Pred. No. 4.9e+04;
Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAAUACGUGAAGA 14
Db 21 AGAAAACATGAGAAGA 34

RESULT 7

```

```

AI039249/c
LOCUS
DEFINITION
Ox33a08.s1 Soares total_fetus.Nb2HF8_9w Homo sapiens cDNA clone
IMAGE:1658102 3' similar to TR:O00554 P21-ARC..[1] ;, mRNA
sequence.
ACCESSION
AI039249
VERSION
AI039249.1 GI:3278443
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1
REFERENCE
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Trace considered overall poor quality
Seq primer: -40ml3 fwd. Et from Amersham
High quality sequence stop: 1.
FEATURES
Location/Qualifiers
1..40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1658102"
/cdb.lib="Soares_total_fetus.Nb2HF8_9w"
/dev_stage="8-9 weeks"
/lab_host="DH10B"
/note="Vector: pT73D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was prepared from mRNA obtained from pooled 8-9 week
(total) fetus material with a Not I - oligo(dT) primer [5',
TGTTACCAATCTGAAGTGGAGCGCGCTTAATTTTTTTTTTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo. "
BASE COUNT 3 a 11 c 10 g 16 t
ORIGIN
Query Match 72.0%; Score 10.8; DB 9; Length 40;
Best Local Similarity 71.4%; Pred. No. 5.1e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAAUACGUGAAGA 14
Db 17 AGATGACGTGATGA 4

RESULT 8
AZ482955
LOCUS
DEFINITION
IM0308M13F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0308M13 F, DNA sequence.
ACCESSION
AZ482955
VERSION
AZ482955.1 GI:10646499
KEYWORDS
GSS.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1
REFERENCE
AUTHORS
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmood,M., Meenen,E., Pedersen,T., Reilly
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D.,Weiss,R.
TITLE
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

```

JOURNAL
COMMENT

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah
Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0308 row: M column: 13
Seq primer: CGTGTAAACGACGCGCCAGT
Class: plasmid ends
High quality sequence stop: 46.
Location/Qualifiers
1. 46

FEATURES
source

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0308M13"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gil147321141gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN

26 a 6 c 8 g 6 t
Query Match 72.0%; Score 10.8; DB 17; Length 46;
Best Local Similarity 78.6%; Pred. No. 5.4e+04;
Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
| | | | | | | | | |
Db 19 AGAAACGTGAAAA 32

RESULT 9
AZ656367

LOCUS 48 bp DNA linear GSS 14-DEC-2000
DEFINITION IM0531D23R Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0531D23 R, DNA sequence.

ACCESSION
AZ656367

VERSION 1 GI:11793513

KEYWORDS
GSS.SOURCE
house mouse.ORGANISM
Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus. 1 (bases 1 to 48)

REFERENCE

1 (bases 1 to 48)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

TITLE

JOURNAL
COMMENT

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah
Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0331 row: D column: 23
Seq primer: CACACAGGAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 48.
Location/Qualifiers
1. 48

FEATURES
source

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0531D23"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gil147321141gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN

26 a 8 c 6 g 8 t
Query Match 72.0%; Score 10.8; DB 17; Length 48;
Best Local Similarity 71.4%; Pred. No. 5.6e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
| | | | | | | | | |
Db 20 AAATAACGTGAAAA 33

RESULT 10
AZ772291

LOCUS 48 bp DNA linear GSS 16-FEB-2001
DEFINITION IM0583L06F Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0583L06 F, DNA sequence.

ACCESSION
AZ772291

VERSION 1 GI:12895445

KEYWORDS
GSS.SOURCE
house mouse.ORGANISM
Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus. 1 (bases 1 to 48)

REFERENCE

1 (bases 1 to 48)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

TITLE

JOURNAL
COMMENT

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0583 row: L column: 06
Seq primer: CGTTGTAACACGCGCCAGT
Class: plasmid ends
High quality sequence stop: 48.

FEATURES

source
1. .48
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0583106"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: pWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (g147321141gb/AP129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 23 a 8 c 7 g 10 t
ORIGIN

Query Match 72.0%; Score 10.8; DB 17; Length 48;
Best Local Similarity 71.4%; Pred. No. 5.6e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGUAACGUGAGA 14
Db 33 AATAACGTGAAAA 46

RESULT 11

AU103231/c
LOCUS AU103231 Sugano Homo sapiens cDNA library EST 30-AUG-2001
DEFINITION KAT07952, mRNA sequence.
ACCESSION AU103231
VERSION AU103231.1 GI:13552752
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 50)
AUTHORS Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J., Hata
Y., Nakamura,Y., Suyama,A. and Sugano,S.
TITLE Diverse transcriptional initiation revealed by fine, large-scale
mapping of mRNA start sites
JOURNAL EMBO Rep. 2 (5), 388-393 (2001)

MEDLINE
COMMENT

21270072
Contact: Yutaka Suzuki
Department of Virology
Institute of Medical Science, University of Tokyo
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
Email: ysuzuki@ims.u-tokyo.ac.jp
Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
S. Construction and characterization of a full length-enriched and
a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES

source
1. .50
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="KAT07952"
/clone_lib="Sugano Homo sapiens cDNA library"
/note="Differential display comparison of untreated and
dimethylfumurate treated U937 cells"
BASE COUNT 17 a 12 c 9 g 12 t
ORIGIN

Query Match 72.0%; Score 10.8; DB 9; Length 50;
Best Local Similarity 71.4%; Pred. No. 5.7e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGAU 15
Db 46 GAGATGTGAAGAT 33

RESULT 12

AVB833384
LOCUS AVB833384 K. Sato unpublished cDNA library: Hordeum vulgare subsp.
DEFINITION vulgare shoots germination Hordeum vulgare subsp. vulgare cDNA
clone bags7123, mRNA sequence.
ACCESSION AVB833384.1 GI:14525473
VERSION AVB833384
KEYWORDS EST.
SOURCE Hordeum vulgare subsp. vulgare.
ORGANISM Hordeum vulgare subsp. vulgare

REFERENCE 1 (bases 1 to 51)
AUTHORS Sato,K.
TITLE Barley EST sequencing project in NIG and Okayama Univ
JOURNAL Unpublished (2001)
COMMENT Contact: Kazuhiro Sato
Research Institute for Bioresources
Okayama University, Barley Germplasm Center
Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
Email: kzsato@rib.okayama-u.ac.jp,
URL:http://www.rib.okayama-u.ac.jp/barley/
Sato,K., Saisho,D., Takeda,K., Shini,T. and Kohara,Y. Direct
submission;
database:http://www.shigen.nig.ac.jp/barley/Barley.html.

FEATURES

source
1. .51
Location/Qualifiers
/organism="Hordeum vulgare subsp. vulgare"
/cultivar="Haruna Nijo"
/db_xref="taxon:112509"
/clone="bags7123"
/clone_lib="K. Sato unpublished cDNA library: Hordeum
vulgare subsp. vulgare shoots germination"
/tissue_type="shoots"
/dev_stage="germination"
BASE COUNT 13 a 14 c 18 g 6 t
ORIGIN

Query Match 72.0%; Score 10.8; DB 10; Length 51;
Best Local Similarity 78.6%; Pred. No. 5.7e+04;
Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

```

Qy      1 AGAUAACGUGAAGA 14
      ||| ||||| |||
Db      36 AGAGAACGCTGGAGA 49

RESULT 13
AA823664
LOCUS   52 bp mRNA linear EST 17-FEB-1998
DEFINITION vt69d09.s1 Knowles Solter mouse 2 cell Mus musculus cDNA clone
IMAGE:1125905 5' similar to gb:J03161 SERUM RESPONSE FACTOR (HUMAN
);, mRNA sequence.
ACCESSION AA823664
VERSION   AA823664.1 GI:2893532
KEYWORDS EST.
SOURCE    house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 52)
Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
Waterston,R.
The WashU-HHMI Mouse EST Project
Unpublished (1996)
Contact: Marra M/Mouse EST Project
WashU-HHMI Mouse EST Project
Washington University School of MedicineP
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:615241.

FEATURES             source
ORIGIN               1..52
                    /organism="Mus musculus"
                    /strain="C57BL/6J x DBA/2J F1"
                    /db_xref="taxon:10090"
                    /clone="IMAGE:1125905"
                    /clone_lib="Knowles Solter mouse 2 cell"
                    /tissue_type="embryo"
                    /dev_stage="2-cell"
                    /lab_host="DH10B"
                    /note="Organ: embryo; Vector: pBluescribe (modified);
                    Site 1: MluI; Site 2: SalI; Cloned unidirectionally from
                    mRNA prepared from 13,500 2-cell stage embryos. Primer:
                    SalI(drr): 5'-CGGTCGACCGTCGACCGTTTCTTTTCTTTT-3'. cDNAs
                    were cloned into the MluI/SalI sites of a modified
                    pBluescribe vector using commercial linkers (NEB).
                    Average insert size: 1.2 kb."
BASE COUNT          26 a 3 c 6 g 17 t
ORIGIN

Query Match      72.0%; Score 10.8; DB 9; Length 52;
Best Local Similarity 71.4%; Pred. No. 5.8e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy      1 AGAUAACGUGAAGA 14
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Db      3 AGATGACGCTGAAAA 16

RESULT 14
BF631905/c
LOCUS   52 bp mRNA linear EST 19-DEC-2000
DEFINITION NF016H07DT1F1061 Drought Medicago truncatula cDNA clone NF016H07DT
5', mRNA sequence.
ACCESSION BF631905
VERSION   BF631905.1 GI:11896063
KEYWORDS EST.

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SOURCE
ORGANISM
barrel medic.
Medicago truncatula
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids 1; Fabales; Fabaceae; Papilionoideae; Trifolieae;
Medicago.
1 (bases 1 to 52)
Torres-Jerez,I., Scott,A.D., Harris,A.R., Gonzales,R.A., Bell,C.J.,
Flores,H.R., Inman,J.T., Wellner,J.W. and May,G.D.
Expressed Sequence tags from the Samuel Roberts Noble Foundation
Medicago truncatula drought library
Unpublished (2000)
Contact: May GD
Plant Biology Division
The Samuel Roberts Noble Foundation
2510 Sam Noble Parkway, Ardmore, OK 73402, USA
Tel: 580 221 7391
Fax: 580 221 7380
Email: gdmay@noble.org
Insert Length: 52 Std Error: 0.00
Plate: 016 Row: H Column: 07
Seq primer: TCACACGAGAAACACGCTATGAC.
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ORIGIN               1..52
                    /organism="Medicago truncatula"
                    /db_xref="taxon:3880"
                    /clone="NF016H07DT"
                    /clone_lib="Drought"
                    /tissue_type="Plantlets"
                    /dev_stage="Pooled timepoints"
                    /note="Vector: Lambda Zap; Contains a mixture of entire
                    plantlets harvested in a series of days-post-watering
                    timepoints."
BASE COUNT          11 a 15 c 5 g 21 t
ORIGIN

Query Match      72.0%; Score 10.8; DB 12; Length 52;
Best Local Similarity 71.4%; Pred. No. 5.8e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy      2 GAUAACGUGAAGAU 15
      ||| ||| |||||
Db      48 GATTAAGAGAGAGAT 35

RESULT 15
AU257573
LOCUS   53 bp mRNA linear EST 25-APR-2002
DEFINITION AU257573 3'-directed mouse cDNA library Mus musculus cDNA clone
BED0010931 3', mRNA sequence.
ACCESSION AU257573
VERSION   AU257573.1 GI:20322326
KEYWORDS EST.
SOURCE    house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 53)
Kato,K. and Matoba,R.
Generation of expressed sequence tags from mouse brain
Unpublished (2002)
Contact: Kikuya Kato
Graduate School of Biological Sciences
Nara Institute of Science and Technology
8916-5 Takayama, Ikoma, Nara 630-0101, Japan
Tel: 81-743-72-5581
Fax: 81-743-72-5589
Email: kkatob@is.aist-nara.ac.jp,
URL:http://love2.aist-nara.ac.jp/BED/index.html.
FEATURES             source
ORIGIN               1..53
                    /organism="Mus musculus"
                    /db_xref="taxon:10090"

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/clone="BED0010931"
/clone_lib="3'-directed mouse cDNA library"
/tissue_type="brain"
/notes="Vector: pGEM-T-easy"

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BASE COUNT      11 a      8 c      20 g      14 t
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Matches 9; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

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Qy      2 GAUACGUGAAGAU 15
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Db      27 GATAGGTCAGAT 40

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Search completed: July 6, 2003, 15:28:26
Job time : 1011.73 secs


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:   OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-5

Query Match          100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 AGAUACGUGAAGAU 15
    |||||
Db  7 AGAUACGUGAAGAU 21

RESULT 8
US-09-780-929-6
: Sequence 6, Application US/09780929
: Patent No. US20020151693A1
: GENERAL INFORMATION:
: APPLICANT: Ribozyme Pharmaceuticals, Inc
: APPLICANT: Breaker, Ronald
: APPLICANT: Beigelman, Leo
: TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
: FILE REFERENCE: MBH00-884-H (500/001)
: CURRENT APPLICATION NUMBER: US/09/780,929
: CURRENT FILING DATE: 2001-02-08
: PRIOR APPLICATION NUMBER: US 60/181,360
: PRIOR FILING DATE: 2000-02-08
: NUMBER OF SEQ ID NOS: 126
: SOFTWARE: PatentIn version 3.0
: SEQ ID NO 6
: LENGTH: 28
: TYPE: RNA
: ORGANISM: Artificial Sequence
: FEATURE:
: OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
: NAME/KEY: misc_feature
: LOCATION: (1)..(4)
: OTHER INFORMATION: 2'-O-Methyl
: NAME/KEY: misc_feature
: LOCATION: (24)..(27)
: OTHER INFORMATION: 2'-O-Methyl
: NAME/KEY: misc_feature
: LOCATION: (28)..(28)
: OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-6

Query Match          100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 AGAUACGUGAAGAU 15
    |||||
Db  7 AGAUACGUGAAGAU 21

RESULT 9
US-09-780-929-7
: Sequence 7, Application US/09780929
: Patent No. US20020151693A1
: GENERAL INFORMATION:
: APPLICANT: Ribozyme Pharmaceuticals, Inc
: APPLICANT: Breaker, Ronald
: APPLICANT: Beigelman, Leo
: TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
: FILE REFERENCE: MBH00-884-H (500/001)
: CURRENT APPLICATION NUMBER: US/09/780,929
: CURRENT FILING DATE: 2001-02-08
: PRIOR APPLICATION NUMBER: US 60/181,360
: PRIOR FILING DATE: 2000-02-08
: NUMBER OF SEQ ID NOS: 126
: SOFTWARE: PatentIn version 3.0
: SEQ ID NO 7
: LENGTH: 28
: TYPE: RNA

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; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(4)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (23)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-7

Query Match          100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 10
US-09-780-929-8
; Sequence 8, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (20)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-8

Query Match          100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 11
US-09-780-929-9
; Sequence 9, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
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; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 9
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (12)..(12)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-9

Query Match          100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 12
US-09-780-929-10
; Sequence 10, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 10
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(7)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-10

Query Match          100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Oy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

RESULT 13

US-09-780-929-11
; Sequence 11, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: 2001-02-08
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 11
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-11

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

RESULT 14

US-09-780-929-12
; Sequence 12, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: 2001-02-08
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 12
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl

; NAME/KEY: misc_feature
; LOCATION: (18)..(18)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-12

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

RESULT 15

US-09-780-929-13
; Sequence 13, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: 2001-02-08
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 13
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (17)..(17)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-13

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

Search completed: July 6, 2003, 16:52:32
Job time : 85 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:40:47 ; Search time 1625.91 Seconds
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Title: US-09-780-929-97

Perfect score: 15

Sequence: 1 agaaacgugaagau 15

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 24791104 seqs, 12571243825 residues

Total number of hits satisfying chosen parameters: 12745074

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Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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77: /cgn2_6/ptodata/1/pna/US6033_COMB.seq.*

78: /cgn2_6/ptodata/1/pna/US6034_COMB.seq.*

79: /cgn2_6/ptodata/1/pna/US6035_COMB.seq.*

80: /cgn2_6/ptodata/1/pna/US6036_COMB.seq.*

81: /cgn2_6/ptodata/1/pna/US6037_COMB.seq.*

82: /cgn2_6/ptodata/1/pna/US6038_COMB.seq.*

83: /cgn2_6/ptodata/1/pna/US6039_COMB.seq.*

84: /cgn2_6/ptodata/1/pna/US6040_COMB.seq.*

85: /cgn2_6/ptodata/1/pna/US6041_COMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	100.0	15	30	US-09-780-929-97
2	15	100.0	27	30	US-09-780-929-3
3	15	100.0	27	30	US-09-780-929-126
4	15	100.0	28	30	US-09-780-929-1
5	15	100.0	28	30	US-09-780-929-2
6	15	100.0	28	30	US-09-780-929-4
7	15	100.0	28	30	US-09-780-929-5
8	15	100.0	28	30	US-09-780-929-6
9	15	100.0	28	30	US-09-780-929-7
10	15	100.0	28	30	US-09-780-929-8
11	15	100.0	28	30	US-09-780-929-9
12	15	100.0	28	30	US-09-780-929-10
13	15	100.0	28	30	US-09-780-929-11
14	15	100.0	28	30	US-09-780-929-12
15	15	100.0	28	30	US-09-780-929-13
16	15	100.0	28	30	US-09-780-929-14
17	15	100.0	28	30	US-09-780-929-15
18	15	100.0	28	30	US-09-780-929-16
19	15	100.0	28	30	US-09-780-929-17
20	15	100.0	28	30	US-09-780-929-18
21	15	100.0	28	30	US-09-780-929-19

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22 15 100.0 28 30 US-09-780-929-20 Sequence 20, Appl
23 15 100.0 28 30 US-09-780-929-21 Sequence 21, Appl
24 15 100.0 28 30 US-09-780-929-22 Sequence 22, Appl
25 15 100.0 28 30 US-09-780-929-23 Sequence 23, Appl
26 15 100.0 28 30 US-09-780-929-24 Sequence 24, Appl
27 15 100.0 28 30 US-09-780-929-25 Sequence 25, Appl
28 15 100.0 28 30 US-09-780-929-26 Sequence 26, Appl
29 15 100.0 28 30 US-09-780-929-27 Sequence 27, Appl
30 15 100.0 28 30 US-09-780-929-28 Sequence 28, Appl
31 15 100.0 28 30 US-09-780-929-29 Sequence 29, Appl
32 15 100.0 28 30 US-09-780-929-30 Sequence 30, Appl
33 15 100.0 28 30 US-09-780-929-31 Sequence 31, Appl
34 15 100.0 28 30 US-09-780-929-32 Sequence 32, Appl
35 15 100.0 28 30 US-09-780-929-33 Sequence 33, Appl
36 15 100.0 28 30 US-09-780-929-34 Sequence 34, Appl
37 15 100.0 28 30 US-09-780-929-35 Sequence 35, Appl
38 15 100.0 28 30 US-09-780-929-36 Sequence 36, Appl
39 15 100.0 28 30 US-09-780-929-37 Sequence 37, Appl
40 15 100.0 28 30 US-09-780-929-38 Sequence 38, Appl
41 15 100.0 28 30 US-09-780-929-39 Sequence 39, Appl
42 15 100.0 28 30 US-09-780-929-40 Sequence 40, Appl
43 15 100.0 28 30 US-09-780-929-41 Sequence 41, Appl
44 15 100.0 28 30 US-09-780-929-42 Sequence 42, Appl
45 15 100.0 28 30 US-09-780-929-43 Sequence 43, Appl
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ALIGNMENTS

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RESULT 1
US-09-780-929-97
; Sequence 97, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 97
; LENGTH: 15
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-929-97
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Query Match 100.0%; Score 15; DB 30; Length 15;
Best Local Similarity 100.0%; Pred. No. 3.8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Oy 1 AGAUACGUGAAGAU 15
Db 1 AGAUACGUGAAGAU 15
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RESULT 2
US-09-780-929-3
; Sequence 3, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
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; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 3
; LENGTH: 27
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-929-3
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Query Match 100.0%; Score 15; DB 30; Length 27;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Oy 1 AGAUACGUGAAGAU 15
Db 7 AGAUACGUGAAGAU 21
|||||
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RESULT 3
US-09-780-929-126
; Sequence 126, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 126
; LENGTH: 27
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc.feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc.feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
US-09-780-929-126
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```
Query Match 100.0%; Score 15; DB 30; Length 27;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
Oy 1 AGAUACGUGAAGAU 15
Db 7 AGAUACGUGAAGAU 21
|||||
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```
RESULT 4
US-09-780-929-1
; Sequence 1, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
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; SEQ ID NO 1
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-1

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 5
US-09-780-929-2
; Sequence 2, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 2
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-2

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 6
US-09-780-929-4
; Sequence 4, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
```

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; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 4
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (22)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-4

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 7
US-09-780-929-5
; Sequence 5, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 5
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (23)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-5

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21
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Db      7 AGAUAACGUGAAGAU 21

RESULT 8
US-09-780-929-6
; Sequence 6, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(4)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (24)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-6

Query Match      100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGAUAACGUGAAGAU 15
        |||||
Db      7 AGAUAACGUGAAGAU 21

RESULT 10
US-09-780-929-8
; Sequence 8, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (20)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-8

Query Match      100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGAUAACGUGAAGAU 15
        |||||
Db      7 AGAUAACGUGAAGAU 21

RESULT 11
US-09-780-929-9
; Sequence 9, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 9
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(4)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (23)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
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```
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (12)..(12)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-9
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 12
US-09-780-929-10
; Sequence 10, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 10
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(7)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-10
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 13
US-09-780-929-11
; Sequence 11, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
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; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 11
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-11
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 14
US-09-780-929-12
; Sequence 12, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 12
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (18)..(18)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-12
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21
```

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Db      7 AGAUACGUGAAGAU 21
|||||
RESULT 15
US-09-780-929-13
; Sequence 13, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 13
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (17)..(17)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-13

Query Match      100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 AGAUACGUGAAGAU 15
|||||
Db      7 AGAUACGUGAAGAU 21

Search completed: July 6, 2003, 16:29:53
Job time : 1625.91 secs
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OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:36:16 ; Search time 591.818 Seconds
(without alignments)
885.154 Million cell updates/sec

Title: US-09-780-929-98
Perfect score: 18
Sequence: 1 aaugccuauccggugcga 18

Scoring table: IDENTITY_NUC
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Total number of hits satisfying chosen parameters: 897812

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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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2	18	100.0	29	6	AX214316	Sequence
3	12.8	71.1	20	6	AX295903	Sequence
4	12.8	71.1	24	6	AX291270	Sequence
5	12.2	67.8	20	6	AX293899	Sequence
6	12.2	67.8	24	6	AX289266	Sequence
7	12.2	67.8	29	6	AR198859	Sequence
8	12	66.7	22	6	AX297915	Sequence
9	11.8	65.6	20	6	AX293092	Sequence
10	11.8	65.6	20	6	E30812	Novel prote
11	11.8	65.6	22	6	AX019596	Sequence
12	11.8	65.6	24	6	AX288459	Sequence
13	11.8	65.6	24	6	AX447444	Sequence
14	11.8	65.6	30	6	A29209	DNA probe f
15	11.8	65.6	30	6	A29212	Oligonucleo
16	11.8	65.6	40	6	AR178716	Sequence
17	11.8	65.6	40	6	AR205421	Sequence
18	11.6	64.4	22	6	AX166857	Sequence
19	11.6	64.4	24	6	AX166856	Sequence
20	11.6	64.4	25	6	AX166855	Sequence
21	11.6	64.4	26	6	I22031	Sequence 5
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27	11.6	64.4	30	6	AR140283	Sequence
28	11.6	64.4	30	6	AR140561	Sequence
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31	11.6	64.4	41	6	AR091476	Sequence
32	11.6	64.4	46	6	AR170886	Sequence
33	11.6	64.4	47	6	AX378262	Sequence
34	11.4	63.3	17	6	AR057682	Sequence
35	11.4	63.3	17	6	AR057773	Sequence
36	11.4	63.3	17	6	AR115440	Sequence
37	11.4	63.3	17	6	AR115531	Sequence
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40	11.4	63.3	24	6	AX447141	Sequence
41	11.4	63.3	25	6	AR097506	Sequence
42	11.4	63.3	25	6	AR139820	Sequence
43	11.4	63.3	25	6	AR140127	Sequence
44	11.4	63.3	25	6	AR142844	Sequence
45	11.4	63.3	31	6	AX425973	Sequence

ALIGNMENTS

RESULT 1
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LOCUS AX214296 18 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 109 from Patent WO0159102.
ACCESSION AX214296
VERSION AX214296.1 GI:15524373
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 18)
AUTHORS Breaker R. and Emilsson G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 109 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)

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RESULT 2
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DEFINITION Sequence 129 from Patent WO0159102.
ACCESSION AX214316
VERSION AX214316.1 GI:15524393
KEYWORDS
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  artificial sequences.
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DEFINITION Sequence 7665 from Patent WO0179548.
ACCESSION AX295903
VERSION AX295903.1 GI:17057592
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  artificial sequences.
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Db 2 ACGGCTTATCGGTGCG 17

RESULT 4
LOCUS AX291270 24 bp DNA linear PAT 21-NOV-2001
DEFINITION Sequence 3032 from Patent WO0179548.
ACCESSION AX291270
VERSION AX291270.1 GI:17052953
KEYWORDS
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  Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

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Db 2 ACGGCTTATCGGTGCG 17

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DEFINITION Sequence 5661 from Patent WO0179548.
ACCESSION AX293899
VERSION AX293899.1 GI:17055582
KEYWORDS
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RESULT 6
LOCUS AX289266 24 bp DNA linear PAT 21-NOV-2001
DEFINITION Sequence 1028 from Patent WO0179548.
ACCESSION AX289266
VERSION AX289266.1 GI:17050949
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Barany,F., Zirvi,M., Gerry,N.P., Favis,R. and Kliman,R.
TITLE Method of designing addressable array for detection of nucleic acid
sequence differences using ligase detection reaction
JOURNAL Patent: WO 0179548-A 1028 25-OCT-2001;
CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES Location/Qualifiers
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Best Local Similarity 64.7%; Pred. No. 5.5e+04;
Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
I:|||||:|||||
Db 7 ATGACCAATCGATGCGA 23

RESULT 7
LOCUS AR198859 29 bp DNA linear PAT 20-APR-2002
DEFINITION Sequence 144 from patent US 6355411.
ACCESSION AR198859
VERSION AR198859.1 GI:20248933
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 29)
AUTHORS Ausubel,F., Goodman,H.M., Rahme,L.G., Mahajan-Miklos,S., Tan,M.-W.,
Cao,H., Drenkard,E. and Tsongalis,J.
TITLE Virulence-associated nucleic acid sequences and uses thereof
JOURNAL Patent: US 6355411-A 144 12-MAR-2002;
FEATURES Location/Qualifiers
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BASE COUNT 6 a 10 c 8 g 5 t
ORIGIN

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Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

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LOCUS AX297915 22 bp DNA linear PAT 26-NOV-2001
DEFINITION Sequence 4 from Patent WO0183757.
ACCESSION AX297915
VERSION AX297915.1 GI:17128036
KEYWORDS TT virus.
SOURCE

ORGANISM TT virus
VIRUSES; ssDNA viruses; unclassified ssDNA viruses.
REFERENCE 1
AUTHORS Ott,C. and Komurian-Pradel,F.
TITLE TT virus polypeptide, nucleic acid coding for said polypeptide and
uses
JOURNAL Patent: WO 0183757-A 4 08-NOV-2001;
BIO MERIEUX (PR)
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DEFINITION Sequence 4854 from Patent WO0179548.
ACCESSION AX293092
VERSION AX293092.1 GI:17054775
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Barany,F., Zirvi,M., Gerry,N.P., Favis,R. and Kliman,R.
TITLE Method of designing addressable array for detection of nucleic acid
sequence differences using ligase detection reaction
JOURNAL Patent: WO 0179548-A 4854 25-OCT-2001;
CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES Location/Qualifiers
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Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

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Db 6 TGCCCTATCTGTGCG 20

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LOCUS E30812/c 20 bp DNA linear PAT 18-JUN-2001
DEFINITION Novel protein participating in differentiation of cranial nerve
tissue cell.
ACCESSION E30812
VERSION E30812.1 GI:13017242
KEYWORDS JP 199318468-A/6.
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Shigeru,N. and Hirofumi,S.
TITLE Novel protein participating in differentiation of cranial nerve
tissue cell
JOURNAL Patent: JP 199318468-A 6 24-NOV-1999;

THE KANAGAWA ACADEMY OF SCIENCE, MEIJI MILK PROD CO LTD
 OS Unidentified
 PN JP 199318468-A/6
 PD 24-NOV-1999
 PF 15-MAY-1998 JP 1998152027
 PR SHIGERU NOGUCHI, HIROFUMI SUEMORI
 PI C12N15/09, A01K67/027, A61K38/00, A61K48/00, C07K14/47, C07K16/18,
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 LOCUS 22 bp DNA linear PAT 07-SEP-2000
 DEFINITION Sequence 50 from Patent WO9938964.
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 ACCESSION AX019596
 VERSION AX019596.1 GI:10043510
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM
 1 (bases 1 to 22)
 REFERENCE
 1 Keith, W.N.
 AUTHORS Keith, W.N.
 TITLE Promoter regions of the mouse and human telomerase rna component
 JOURNAL
 Patent: WO 9938964-A 50 05-AUG-1999;
 KEITH WILLIAM NICOL (GB); CANCER RES CAMPAIGN TECH (GB)
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 LOCUS 24 bp DNA linear PAT 21-NOV-2001
 DEFINITION Sequence 221 from Patent WO0179548.
 AX288459
 ACCESSION AX288459
 VERSION AX288459.1 GI:17050142
 KEYWORDS

synthetic construct.
 ORGANISM
 synthetic construct
 artificial sequences.
 REFERENCE
 1 Barany, F., Zivvi, M., Gerry, N.P., Favis, R. and Kliman, R.
 AUTHORS Method of designing addressable array for detection of nucleic acid
 TITLE sequence differences using ligase detection reaction
 JOURNAL Patent: WO 0179548-A 221 25-OCT-2001;
 CORNELL RESEARCH FOUNDATION, INC. (US)
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 6 TGGCTATCTGTGGC 20
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 AX447444
 LOCUS 24 bp DNA linear PAT 03-JUL-2002
 DEFINITION Sequence 3899 from Patent WO0216649.
 AX447444
 ACCESSION AX447444
 VERSION AX447444.1 GI:21696343
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM
 1 synthetic construct
 artificial sequences.
 REFERENCE
 1 Gunderson, K.
 AUTHORS Probes and decoder oligonucleotides
 TITLE Patent: WO 0216649-A 3899 28-FEB-2002;
 JOURNAL Illumina, Inc. (US)
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 RESULT 14
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 LOCUS 30 bp DNA linear PAT 30-JUN-1995
 DEFINITION DNA probe from patent WO9111459.
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 ACCESSION AX29209
 VERSION AX29209.1 GI:1248930
 KEYWORDS
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 AUTHORS
 TITLE
 JOURNAL
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GenCore version 5.1.6
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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6	12.8	71.1	24	24 ABI88460	Capture oligonucle
7	12.8	71.1	24	24 ABI88461	Capture oligonucle
8	12.4	68.9	41	22 AAC87378	Staphylococcus aur
9	12.2	67.8	20	24 ABI93941	Capture oligonucle

c 10	12.2	67.8	24	22 AAS08711	Human PD-ATP-bindin
c 11	12.2	67.8	24	24 ABI84452	Capture oligonucle
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c 13	12.2	67.8	29	20 AAX98255	PCR primer used to
c 14	12.2	67.8	40	17 AAT70787	Stenotic carotid a
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c 16	12.2	67.8	60	24 ABN32869	Human spliced tran
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c 18	11.8	65.6	20	20 AAX94010	PCR primer used to
c 19	11.8	65.6	20	21 AAZ44197	Murine cerebral ne
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c 30	11.8	65.6	40	21 AAA62656	Beta-lactamase gen
c 31	11.6	64.4	22	22 AAD11273	Mycobacterium 16S
c 32	11.6	64.4	24	22 AAD11272	Mycobacterium 16S
c 33	11.6	64.4	25	22 AAD11271	Mycobacterium 16S
c 34	11.6	64.4	27	19 AAV55972	Human cytokine rec
c 35	11.6	64.4	29	16 AAQ93711	Primer corresp. to
c 36	11.6	64.4	29	17 AAT10032	MDGF antisense pri
c 37	11.6	64.4	30	16 AAT04920	Mammalian stem cel
c 38	11.6	64.4	30	16 AAQ99710	Primer for reverse
c 39	11.6	64.4	30	17 AAT10031	MDGF cDNA/antisens
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ALIGNMENTS

RESULT 1
AAS12348
ID AAS12348 standard; DNA; 18 BP.

XX AAS12348;

XX 21-NOV-2001 (first entry)

XX DNA encoding deoxyribozyme #8.

XX Deoxyribozyme; cytosstatic; endonuclease; RNA cleavage; DNA cleavage;

XX gene therapy; plant; fungus; bacteria; mammal; ribozyme; ss.

XX Synthetic.

XX WO200159102-A2.

XX 16-AUG-2001.

XX 08-FEB-2001; 2001WO-US04223.

XX 08-FEB-2000; 2000US-0181360.

XX 31-MAR-2000; 2000US-0193646.

XX (RIBO-) RIBOZYME PHARM INC.

XX (UYIA) UNIV YALE.

XX Breaker R, Belgelman L, Emilsson G;

XX WPI; 2001-536526/59.

XX New nucleic acids with endonuclease activity, such as ribozymes and

PT nucleozymes, for modulating gene expression in a plant, mammalian,
 PT bacterial or fungal cell -
 PS Claim 49; Page 77; 96pp; English.
 XX
 CC The invention relates to nucleic acid molecules with endonuclease
 CC activity, which are particularly useful for cleavage of RNA or DNA.
 CC The nucleic acids are used in a pharmaceutical composition and are used
 CC to modulate expression of a gene in a plant, mammalian, bacterial or
 CC fungal cell. They are used to cleave a separate nucleic acid, preferably
 CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
 CC proliferation, and can be used to treat a disease or condition. More
 CC than one nucleic acid can be independently targeted to the same or
 CC different sites in a cell. The nucleic acids may be used to study DNA.
 CC The modifications to the nucleic acids optimises their catalytic activity
 CC and can maintain or enhance their activity. They exhibit a high degree
 CC of specificity for RNA. The present sequence represents the coding
 CC sequence of deoxyribozyme #8 used in the method of the invention.
 XX
 XX Sequence 18 BP; 4 A; 4 C; 6 G; 4 U; 0 other;

Query Match 100.0%; Score 18; DB 22; Length 18;
 Best Local Similarity 100.0%; Pred. No. 0.63;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AAUGGCCUAUCGGUGCGA 18
 |||||
 Db 1 AAUGGCCUAUCGGUGCGA 18

RESULT 2
 AAS12381
 ID AAS12381 standard; RNA; 29 BP.
 XX
 AC AAS12381;
 XX
 DT 21-NOV-2001 (first entry)
 XX
 DE Class IV ribozyme.
 XX
 KW Deoxyribozyme; cytotatic; endonuclease; RNA cleavage; DNA cleavage;
 KW gene therapy; plant; fungus; bacteria; mammal; ribozyme; ss.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT misc_binding 1..8
 FT /*tag= a
 FT /note= "Forms double-stranded region with bases 15
 FT to 8 of AAS12374"
 FT
 FT misc_binding 25..29
 FT /*tag= b
 FT /note= "Forms double-stranded region with bases 5
 FT to 1 of AAS12374"
 FT
 XX WO200159102-A2.
 XX
 PD 16-AUG-2001.
 XX
 PF 08-FEB-2001; 2001WO-US04223.
 XX
 XX 08-FEB-2000; 2000US-0181360.
 PR 31-MAR-2000; 2000US-0193646.
 XX
 XX (RIBO-) RIBOZYME PHARM INC.
 PA (UYVA) UNIV YALE.
 PA
 PI Breaker R, Beigelman L, Emilsson G;
 XX
 XX WPI; 2001-536526/59.
 DR
 XX New nucleic acids with endonuclease activity, such as ribozymes and
 PT nucleozymes, for modulating gene expression in a plant, mammalian,
 PT

PT bacterial or fungal cell -
 XX
 PS Example 1; Fig 9; 96pp; English.
 XX
 CC The invention relates to nucleic acid molecules with endonuclease
 CC activity, which are particularly useful for cleavage of RNA or DNA.
 CC The nucleic acids are used in a pharmaceutical composition and are used
 CC to modulate expression of a gene in a plant, mammalian, bacterial or
 CC fungal cell. They are used to cleave a separate nucleic acid, preferably
 CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
 CC proliferation, and can be used to treat a disease or condition. More
 CC than one nucleic acid can be independently targeted to the same or
 CC different sites in a cell. The nucleic acids may be used to study DNA.
 CC The modifications to the nucleic acids optimises their catalytic activity
 CC and can maintain or enhance their activity. They exhibit a high degree
 CC of specificity for RNA. The present sequence represents the Class IV
 CC ribozyme, used in an example which demonstrates the method of
 CC the invention.
 XX
 XX Sequence 29 BP; 6 A; 7 C; 11 G; 5 U; 0 other;

Query Match 100.0%; Score 18; DB 22; Length 29;
 Best Local Similarity 100.0%; Pred. No. 0.67;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AAUGGCCUAUCGGUGCGA 18
 |||||
 Db 8 AAUGGCCUAUCGGUGCGA 25

RESULT 3
 AAS19241/c
 ID AAS19241 standard; DNA; 30 BP.
 XX
 AC AAS19241;
 XX
 DT 26-MAR-2002 (first entry)
 XX
 DE CMV mutagenic adaptor.
 XX
 KW T0; ds; terminator; pGA; DNA vaccine; anti-HIV; virucide;
 KW Human Immunodeficiency Virus; HIV; Gag; HIV gp120; HIV Env;
 KW HIV VLP; measles fusion protein; measles haemagglutinin; CMV; adaptor;
 KW measles nucleoprotein; influenza haemagglutinin; C3d gene;
 KW cell-mediated immune response; humoral immune response; infection.
 XX
 OS Human cytomegalovirus.
 XX
 PN WO200192470-A2.
 XX
 PD 06-DEC-2001.
 XX
 PF 02-MAR-2001; 2001WO-US06795.
 XX
 PR 02-MAR-2000; 2000US-186364P.
 PR 01-DEC-2000; 2000US-251083P.
 XX
 PA (UYEM-) UNIV EMORY.
 XX
 PI Robinson HL, Smith JM, Ross TM, Bright RA, Hua J, Ellenberger D;
 XX
 DR WPI; 2002-075465/10.
 XX
 PT Novel pGA vector useful for immunising patient against measles,
 PT influenza has termination sequence encoding lambda T0 terminator and a
 PT eukaryotic transcription cassette with vaccine insert encoding
 PT immunogens of pathogens -
 XX
 XX Example 2; Page 43; 174pp; English.
 PS
 CC The invention relates to a vector (a pGA construct) comprising a
 CC termination sequence coding for the lambda T0 terminator, a prokaryotic
 CC origin of replication, a selectable marker gene and a eukaryotic

transcription cassette comprising a vaccine insert encoding one or more immunogens derived from a pathogen e.g. Human Immunodeficiency Virus (HIV) Gag, HIV gp120, HIV Pol, HIV Env, HIV VLP, or its mutants, measles fusion protein, measles haemagglutinin, measles nucleoprotein, influenza haemagglutinin, or its mutants, or subsequences, and optionally at least one C3d gene, is useful for immunising or treating a patient, when administered by an intramuscular or intradermal route. The immunisation methods using pGA elicit both cell-mediated and humoral immune responses that may limit the infection, spread or growth of the pathogen and result in protection against subsequent challenge against the pathogen. The terminator sequence present prevents read-through from the kanamycin cassette into vaccine sequences while the plasmid is being produced in bacteria. Prevention of transcriptional read-through stabilises vaccine insert sequences by limiting the exposure of secondary structures that can be recognised by bacterial endonucleases. The present sequence is an adaptor for introducing a ClaI site into the cytomegalovirus (CMV) promoter of pGA2.

XX SQ Sequence 30 BP; 7 A; 12 C; 6 G; 5 T; 0 other;

Query Match 76.7%; Score 13.8; DB 24; Length 30;
Best Local Similarity 64.7%; Pred. No. 1.9e+02;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
I:||||:|||||
Db 25 ATGGCGTATCGATCGGA 9

RESULT 4

ABN47743
ID ABN47743 standard; DNA; 60 BP.

XX AC ABN47743;

XX DT 15-JUL-2002 (first entry)

XX DE Human spliced transcript detection oligonucleotide SEQ ID NO:20491.

XX KW Human; mouse; rat; splice transcript; detection; RNA transcript;
KW splice variant; transcriptome; oligonucleotide library; ss.

XX OS Homo sapiens.

XX PN WO200210449-A2.

XX PD 07-FEB-2002.

XX PF 20-JUL-2001; 2001WO-IB01903.

XX PR 28-JUL-2000; 2000US-221607P.

XX PR 02-MAY-2001; 2001US-287724P.

XX PA (COMP-) COMPUGEN INC.

XX PI Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX DR WPI; 2002-257383/30.

XX PT New oligonucleotide libraries comprising oligonucleotides which
PT selectively hybridize to mRNAs transcribed from a transcription unit of
PT a genome, useful for detecting tissue-, pathology-, and
PT developmental-specific genes -

XX PS Example 1; SEQ ID 20491; 47pp; English.

XX CC The present invention describes oligonucleotide libraries for detecting
CC messenger RNAs that populate a (sub-)transcriptome, where the
CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple
CC transcription units that populate a genome. The library comprises
CC several oligonucleotides, each capable of hybridising selectively to a
CC set of messenger RNAs transcribed from a given transcription unit of
CC the genome, which encodes one or more messenger RNA splice variants.

CC The oligonucleotide libraries are useful for detecting mRNAs from a
CC biological sample, in expression profiling studies, in qualitatively or
CC quantitatively characterising the corresponding transcriptome, and in
CC detecting RNA transcripts and splice variants of human or animal
CC transcriptomes. The libraries may also be used as specialised mini
CC libraries to detect transcripts of a sub-transcriptome under a
CC particular biological or pathological state, and so allowing the
CC detection of tissue- and pathology-specific genes such as those genes
CC only expressed in specific tissue under a specific pathological
CC condition; to detect developmental specific genes; and to detect RNA
CC transcripts and splice variants of a transcriptome of a patient suffering
CC from a particular disorder. ABN27253 to ABN59589 represent
CC oligonucleotide sequences from rats, humans and mice, which are used in
CC the exemplification of the present invention.

CC N.B. The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 60 BP; 16 A; 18 C; 13 G; 13 T; 0 other;

Query Match 74.4%; Score 13.4; DB 24; Length 60;
Best Local Similarity 66.7%; Pred. No. 3.6e+02;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
I:||||:|||||
Db 1 ATGGCGTATCGGTG 15

RESULT 5

ABI95945
ID ABI95945 standard; DNA; 20 BP.

XX AC ABI95945;

XX DT 16-FEB-2002 (first entry)

XX DE Capture oligonucleotide Zip ID#3032 oligo #9.

XX KW Human; K-ras; PCR primer; probe; capture probe; mutation detection;
KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
KW environmental monitoring; food industry; feed industry; ss.

XX OS Synthetic.

XX PN WO200179548-A2.

XX PD 25-OCT-2001.

XX PF 04-APR-2001; 2001WO-US10958.

XX PR 14-APR-2000; 2000US-197271P.

XX PA (CORR) CORNELL RES FOUND INC.

XX PI Barany F, Zirvi M, Gerry NP, Favis R, Kilman R;

XX DR WPI; 2002-034366/04.

XX PT Designing capture oligonucleotide probes for use on a support to which
PT complementary oligonucleotides hybridize with little mismatch -

XX PS Example 5; Fig 29; 300pp; English.

XX CC The present invention describes a method (M1) for designing capture
CC oligonucleotide probes (I) for use on a support to which complementary
CC oligonucleotide probes (II) will hybridise with little mismatch, where
CC (I) have melting temperatures within a narrow range. The method is useful
CC for detecting infectious diseases caused by bacterial infectious agents
CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and

CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. AB182074 to
 CC AB197546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX
 SQ Sequence 20 BP; 2 A; 6 C; 6 G; 6 T; 0 other;

Query Match 71.1%; Score 12.8; DB 24; Length 20;
 Best Local Similarity 68.8%; Pred. No. 7.2e+02;
 Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUGGUGCG 17
 1 111 :1:111:111
 Db 2 ACGGCTTATCGGTGCG 17

RESULT 6

AB188460
 ID AB188460 standard; DNA; 24 BP.

AC AB188460;

XX
 DT 15-FEB-2002 (first entry)

DE Capture oligonucleotide Zip ID#3032 oligo #1.

XX Human; K-ras; PCR primer; probe; capture probe; mutation detection;
 KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
 KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
 KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
 KW environmental monitoring; food industry; feed industry; ss.

OS Synthetic.

XX WO200179548-A2.

XX 25-OCT-2001.

XX 04-APR-2001; 2001WO-US10958.

XX 14-APR-2000; 2000US-197271P.

XX (CORR) CORNELL RES FOUND INC.

XX Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;

XX WPI; 2002-034366/04.

XX Designing capture oligonucleotide probes for use on a support to which
 XX complementary oligonucleotides hybridize with little mismatch -

XX Example 5; Fig 25; 300pp; English.

XX The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and

CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. AB182074 to
 CC AB197546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX
 SQ Sequence 24 BP; 2 A; 7 C; 7 G; 8 T; 0 other;

Query Match 71.1%; Score 12.8; DB 24; Length 24;
 Best Local Similarity 68.8%; Pred. No. 7.3e+02;
 Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUGGUGCG 17
 1 111 :1:111:111
 Db 2 ACGGCTTATCGGTGCG 17

RESULT 7

AB188461/G

ID AB188461 standard; DNA; 24 BP.

XX
 AC AB188461;

XX
 DT 15-FEB-2002 (first entry)

DE Capture oligonucleotide Zip ID#3032 oligo #2.

XX Human; K-ras; PCR primer; probe; capture probe; mutation detection;
 KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
 KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
 KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
 KW environmental monitoring; food industry; feed industry; ss.

OS Synthetic.

XX WO200179548-A2.

XX 25-OCT-2001.

XX 04-APR-2001; 2001WO-US10958.

XX 14-APR-2000; 2000US-197271P.

XX (CORR) CORNELL RES FOUND INC.

XX Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;

XX WPI; 2002-034366/04.

XX Designing capture oligonucleotide probes for use on a support to which
 XX complementary oligonucleotides hybridize with little mismatch -

XX Example 5; Fig 25; 300pp; English.

XX The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and

Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus, Epstein-Barr virus, and parasitic infectious agents selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus medinensis. The method is also useful for detecting genetic diseases such as 21 hydroxylase deficiency, Turner Syndrome and obesity defects. Detecting cancer involving oncogenes, tumour suppressor genes, or genes involved in DNA amplification, replication, recombination or repair, the p53 gene, human papillomavirus types 16 and 18 and liver cancers. The method is also used for environmental monitoring, forensics and the food and feed industry, detecting comprises scanning (using e.g. a scanning electron microscope and infrared microscope) the support at the particular sites and identifying if ligation of the oligonucleotide probe sets occurred and correlating (using a computer) identified ligation to a presence or absence of the target nucleotide sequences. ABI82074 to ABI97546 represent oligonucleotide sequences used in the exemplification of the present invention.

Sequence 24 BP; 8 A; 7 C; 7 G; 2 T; 0 other;

Query Match 71.1%; Score 12.8; DB 24; Length 24;
Best Local Similarity 68.8%; Pred. No. 7.3e+02;
Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AUGGCCUAUCGGUGCG 17
| | | | : | : | | | | |
Db 23 ACGGCTTATCGGTGCG 8

RESULT 8
AAC87378
ID AAC87378 standard; DNA; 41 BP.

XX AAC87378;

XX 09-MAR-2001 (first entry)

XX Staphylococcus aureus Spa domain D antisense PCR primer, PC3H AS.

XX Spa domain D; randomised library; VH3 Ig-Fab fragment; immunoglobulin; Spa mutant; superantigen; altered specificity; apoptosis inducer; energy inducer; B-lymphocyte subset; B-cell; lymphoma; leukaemia; autoimmune disease; idiopathic thrombocytopenia; rheumatoid arthritis; systemic lupus erythematosus; SLE; autoimmune thyroiditis; diabetes; antibody purification; PCR primer; ss.

XX Staphylococcus aureus.

XX WO2000069457-A1.

XX 23-NOV-2000.

XX 15-MAY-2000; 2000WO-US13402.

XX 15-MAY-1999; 99US-0134386.

XX (UYCA-) UNIV CALIFORNIA SAN DIEGO.

XX Silverman GJ;

XX WPI; 2001-031886/04.

XX New staphylococcal protein A variant, useful for treating diabetes and rheumatoid arthritis, exhibits binding specificity for immunoglobulin-Fab domain and comprises variations in amino acid sequence of staphylococcal protein A domain D.

XX Example 12; Page 58; 88pp; English.

XX The invention relates to staphylococcal protein A (SpA) variants which exhibit altered binding specificity for an immunoglobulin Fab (Ig-Fab) fragment relative to native SpA. The SpA variants of the invention have one or more amino acid substitution in the SpA VH3 Ig-Fab binding region

(i.e., SpA domain D) relative to the native SpA. The SpA variants are inducers of autoreactive B cell, leukaemic or lymphoma cell apoptosis or energy. The SpA variants of the invention are useful for detecting the presence of a certain Ig-Fab-expressing lymphocyte subset in a sample of lymphocytes. The SpA variants may therefore be used in the diagnosis of some forms of leukaemia. The SpA variants may also be administered to an individual with an abnormally high number of a certain lymphocyte subset to reduce the number of that lymphocyte subset. The SpA variants are also useful for purifying monoclonal or polyclonal antibodies from serum, plasma, tissue culture or other sources. The SpA variants that exhibit enhanced clon VH3-specific effects in vitro or in vivo, such as the ability to delete undesirable neoplastic B-cells or pathogenic B-cells that are responsible for the production of disease-causing autoantibodies, are useful as therapeutic agents. Therapeutic SpA variants that can bind to Fab on the B-cell receptor of an autoreactive B-cell or leukaemic or lymphoma cell can induce energy, apoptosis, or deletion by other mechanisms. By engineering the interaction between SpA variants and Ig-Fab according to the invention, variant SpA with specially tailored Fab-binding specificities can be selected that target pathogenic neoplastic B cell populations or autoreactive B-cell clones. The SpA variants are therefore useful for treating conditions that are linked to disease-associated B-cells such as idiopathic thrombocytopenia, rheumatoid arthritis, systemic lupus erythematosus (SLE), autoimmune thyroiditis, or diabetes. Engineered B cell superantigens (such as the SpA variants of the invention) bind to immunoglobulin receptors on B-cells in a manner that is distinct from antigen binding by antibodies. Therefore, the SpA variants which have superantigen properties can target much larger populations of Ig-expressing B cells in vivo compared with targeting using a specific antigen. Sequences AAC87372-C87378 represent nucleic acid sequences used in the construction of a randomised SpA domain D library in an exemplification of the invention. Sequences AAC87375-C87378 represent SpA domain D PCR primers used in library construction.

SQ Sequence 41 BP; 5 A; 9 C; 19 G; 8 T; 0 other;

Query Match 68.9%; Score 12.4; DB 22; Length 41;
Best Local Similarity 64.3%; Pred. No. 1.3e+03;
Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 3 UGGCCUAUCGGUGCG 16
| | | | : | : | | | | |
Db 23 TGGCCTTTCGGTGC 36

RESULT 9

ABI93941

ID ABI93941 standard; DNA; 20 BP.

XX ABI93941;

XX 16-FEB-2002 (first entry)

XX Capture oligonucleotide Zip ID#1028 oligo #9.

XX Human; K-ras; PCR primer; probe; capture probe; mutation detection; ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease; infection; 21 hydroxylase deficiency; Turner Syndrome; obesity; cancer; oncogene; tumour suppressor; human papillomavirus; forensic; environmental monitoring; food industry; feed industry; ss.

XX Synthetic.

XX WO200179548-A2.

XX 25-OCT-2001.

XX 04-APR-2001; 2001WO-US10958.

XX 14-APR-2000; 2000US-197271P.

XX (CORR) CORNELL RES FOUND INC.

XX

PI Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;
XX WPI; 2002-034366/04.
XX
PT Designing capture oligonucleotide probes for use on a support to which
PT complementary oligonucleotides hybridize with little mismatch -
XX
PS Example 5; Fig 29; 300pp; English.
XX
CC The present invention describes a method (M1) for designing capture
CC oligonucleotide probes (I) for use on a support to which complementary
CC oligonucleotide probes (II) will hybridize with little mismatch, where
CC (I) have melting temperatures within a narrow range. The method is useful
CC for detecting infectious diseases caused by bacterial infectious agents
CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenzae, fungal
CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and
CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
CC Epstein-Barr virus and polio virus, and parasitic infectious agents
CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
CC medinensis. The method is also useful for detecting genetic diseases such
CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
CC involved in DNA amplification, replication, recombination or repair, the
CC cancer is specifically associated with a gene selected from BRCA1 gene,
CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
CC method is also used for environmental monitoring, forensics and the food
CC and feed industry, detecting comprises scanning (using e.g. a scanning
CC electron microscope and infrared microscope) the support at the
CC particular sites and identifying if ligation of the oligonucleotide probe
CC sets occurred and correlating (using a computer) identified ligation to a
CC presence or absence of the target nucleotide sequences. ABI82074 to
CC ABI97546 represent oligonucleotide sequences used in the exemplification
CC of the present invention.
XX
SQ Sequence 20 BP; 7 A; 6 C; 4 G; 3 T; 0 other;

Query Match 67.8%; Score 12.2; DB 24; Length 20;
Best Local Similarity 64.7%; Pred. No. 1.6e+03;
Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
Qy 2 AUGGCCUAUCGGUGCGA 18
I : I I I I I I I I I I
Db 3 ATGACCAATCGATCGCA 19
RESULT 10
AAS08711/c
ID AAS08711 standard; DNA; 24 BP.
AC AAS08711;
XX
DT 26-SEP-2001 (first entry)
XX
DE Human PD-ATP-binding cassette (PD-ABC) cDNA reverse PCR primer #2.
XX
KW PD-ATP-binding cassette; PD-ABC; chromosome 19p13.3; spleen; thymus; ss;
KW peripheral blood leukocyte; bone marrow; lymph node; dyslipidaemia;
KW cardiovascular disorder; inflammatory disorder; abnormal calcium flux;
KW epilepsy; coronary artery disease; Tangier's disease; atherosclerosis;
KW familial high-density lipoprotein deficiency; fatty liver disease;
KW atherosclerosis; diabetes; insulin resistance; obesity; drug screening;
KW alcoholism; retinal degeneration; hypertension; vascular disease;
KW PCR primer.
XX
OS Synthetic.
XX
PN WO200153490-A1.
XX
PD 26-JUL-2001.
XX
PF 23-JAN-2001; 2001WO-US02191.
XX
PS 24-JAN-2000; 2000US-0177899.

PR 30-JUN-2000; 2000US-0215405.
XX (WARN) WARNER LAMBERT CO.
XX
PI Johns MA, Tafuri SR, Wang M;
XX WPI; 2001-442259/47.
XX
PT New Human PD-ABC DNA molecules and proteins for diagnosis and treatment
PT of dyslipidaemia, epilepsy and diseases related to abnormal calcium flux
XX
PS Disclosure; Page 34; 77pp; English.
XX
CC The sequence represents a PCR primer used for isolation of cDNA encoding
CC human PD-ATP-binding cassette (PD-ABC) protein. PD-ABC maps to chromosome
CC 19p13.3 and is expressed in various tissues including spleen, thymus,
CC peripheral blood leukocytes, bone marrow and lymph nodes. The PD-ABC DNA
CC molecules and proteins are used to diagnose and treat cardiovascular
CC disorders, inflammatory disorders, dyslipidaemia, epilepsy, diseases
CC related to abnormal calcium flux, coronary artery disease, Tangier's
CC disease, familial high-density lipoprotein deficiency, atherosclerosis,
CC diabetes, fatty liver disease, insulin resistance, obesity, alcoholism,
CC retinal degeneration, hypertension and vascular disease. The sequences
CC are also used in drug screening assays.
XX
SQ Sequence 24 BP; 7 A; 6 C; 7 G; 4 T; 0 other;
Query Match 67.8%; Score 12.2; DB 22; Length 24;
Best Local Similarity 58.8%; Pred. No. 1.6e+03;
Matches 10; Conservative 4; Mismatches 3; Indels 0; Gaps 0;
Qy 2 AUGGCCUAUCGGUGCGA 18
I : I I I I I I I I I I
Db 24 ATGCCCTATCCGTGCTA 8
RESULT 11
ABI84452
ID ABI84452 standard; DNA; 24 BP.
AC ABI84452;
XX
DT 15-FEB-2002 (first entry)
XX
DE Capture oligonucleotide zip ID#1028 oligo #1.
XX
KW Human; K-ras; PCR primer; probe; capture probe; mutation detection;
KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
KW environmental monitoring; food industry; feed industry; ss.
XX
OS Synthetic.
XX
PN WO200179548-A2.
XX
PD 25-OCT-2001.
XX
PF 04-APR-2001; 2001WO-US10958.
XX
PR 14-APR-2000; 2000US-197271P.
XX
PA (CORR) CORNELL RES FOUND INC.
XX
PI Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;
XX WPI; 2002-034366/04.
XX
PT Designing capture oligonucleotide probes for use on a support to which
PT complementary oligonucleotides hybridize with little mismatch -
XX
PS Example 5; Fig 25; 300pp; English.

XX The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and
 CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. ABI82074 to
 CC ABI97546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX SQ Sequence 24 BP; 7 A; 7 C; 5 G; 5 T; 0 other;

Query Match 67.8%; Score 12.2; DB 24; Length 24;
 Best Local Similarity 64.7%; Pred. No. 1.6e+03;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
 I: | | | | |
 Db 7 ATGACCAATCGATCGCA 23

RESULT 12

ABI84453/c
 ID ABI84453 standard; DNA; 24 BP.

XX AC ABI84453;

XX DT 15-FEB-2002 (first entry)

XX DE Capture oligonucleotide zip ID#1028 oligo #2.

XX KW Human; K-ras; PCR primer; probe: capture probe: mutation detection;
 KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
 KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
 KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
 KW environmental monitoring; food industry; feed industry; ss.

XX OS Synthetic.

XX PN WO200179548-A2.

XX PD 25-OCT-2001.

XX PF 04-APR-2001; 2001WO-US10958.

XX PR 14-APR-2000; 2000US-197271P.

XX PA (CORR) CORNELL RES FOUND INC.

XX PI Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;

XX DR WPI; 2002-034366/04.

XX PT Designing capture oligonucleotide probes for use on a support to which
 PT complementary oligonucleotides hybridize with little mismatch -

XX PS Example 5; Fig 25; 300pp; English.

XX

CC The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and
 CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. ABI82074 to
 CC ABI97546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX SQ Sequence 24 BP; 5 A; 5 C; 7 G; 7 T; 0 other;

Query Match 67.8%; Score 12.2; DB 24; Length 24;
 Best Local Similarity 64.7%; Pred. No. 1.6e+03;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
 I: | | | | |
 Db 18 ATGACCAATCGATCGCA 2

RESULT 13

AAAX98255/c
 ID AAAX98255 standard; DNA; 29 BP.

XX AC AAAX98255;

XX DT 25-OCT-1999 (first entry)

XX DE PCR primer used to amplify a 1659 bp fragment containing ORF1.

XX KW Human pathogen; virulence polypeptide; virulence factor;
 KW pathogenic infection; Pseudomonas aeruginosa infection; PCR primer; ss.

XX OS Synthetic.

XX OS Pseudomonas aeruginosa.

XX PN WO927129-A1.

XX PD 03-JUN-1999.

XX PF 25-NOV-1998; 98WO-US25247.

XX PR 25-NOV-1997; 97US-0066517.

XX PA (GEHO) GEN HOSPITAL CORP.

XX PI Ausubel F, Cao H, Drenkard E, Goodman HM, Mahajan-Miklos S;

XX PI Rahme IG, Tan M, Tsongalis J;

XX DR WPI; 1999-357851/30.

XX PT Virulence factors useful in developing disease treatments

XX PS Disclosure; Page 26; 228pp; English.

XX CC The present sequence represents a PCR primer used to amplify Pseudomonas

CC 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
CC untranslated sequence is unique to a particular mRNA species, almost
CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
CC is constructed so as to reflect accurately the relative abundance of
CC different mRNAs in the particular tissue from which it was derived.
CC The appearance frequency of a given GS in a cDNA library can be
CC determined (esp. using primers and probes derived from the GS
CC sequences) as a means of diagnosing abnormal cell function or for
CC recognising different cell types.
XX
SQ Sequence 53 BP; 14 A; 11 C; 12 G; 15 T; 1 other;
Query Match 67.8%; Score 12.2; DB 16; Length 53;
Best Local Similarity 64.7%; Pred. No. 1.8e+03;
Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
Oy 1 AAUGGCCCUAUCGUGCG 17
 ||| :| :| :
Db 34 AATGCCCCCTCGATCG 18

Search completed: July 6, 2003, 14:32:53
Job time : 194.091 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:26:51 ; Search time 1209.27 Seconds
(without alignments)
241.069 Million cell updates/sec

Title: US-09-780-929-98

Perfect score: 18
Sequence: 1 aauggccuauccgugcgga 18

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 146654

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

EST.*
1: em_estba.*
2: em_esthum.*
3: em_estin.*
4: em_estnu.*
5: em_estov.*
6: em_estpl.*
7: em_estro.*
8: em_hic.*
9: gb_est1.*
10: gb_est2.*
11: gb_hic.*
12: gb_est3.*
13: gb_est4.*
14: gb_est5.*
15: em_estfun.*
16: em_estom.*
17: gb_gss.*
18: em_gss_hum.*
19: em_gss_inv.*
20: em_gss_pln.*
21: em_gss_vrt.*
22: em_gss_fun.*
23: em_gss_man.*
24: em_gss_mus.*
25: em_gss_other.*
26: em_gss_pro.*
27: em_gss_rod.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	13.8	76.7	49	17	TA58C07Q
c 2	12.4	68.9	50	9	AU183461
3	12.2	67.8	37	14	H13124
c 4	12.2	67.8	50	14	C00960
5	11.8	65.6	50	9	AU105634
c 6	11.8	65.6	56	10	AV847604

7	11.6	64.4	29	17	AZ800642
8	11.6	64.4	55	9	AA863171
c 9	11.2	62.2	30	17	AZ840293
c 10	11.2	62.2	36	17	AZ605771
11	11.2	62.2	37	9	AA931624
c 12	11.2	62.2	45	17	AZ317769
13	11.2	62.2	51	17	BH618002
14	11.2	62.2	52	17	AA721034
15	11.2	62.2	54	17	BH641243
c 16	11	61.1	32	17	TA371C10P
c 17	11	61.1	58	9	A1221514
c 18	10.8	60.0	32	13	BJ066180
c 19	10.8	60.0	44	17	BH252563
c 20	10.8	60.0	58	9	A1953654
21	10.8	60.0	59	17	AL758565
c 22	10.6	58.9	27	17	TA208E12P
23	10.6	58.9	38	17	TA244G07P
c 24	10.6	58.9	40	17	A1613042
c 25	10.6	58.9	40	17	AZ511081
c 26	10.6	58.9	44	17	AZ495581
c 27	10.6	58.9	46	9	AA455514
c 28	10.6	58.9	50	9	AU102871
c 29	10.6	58.9	50	17	AZ777046
c 30	10.6	58.9	55	17	BH810903
31	10.6	58.9	56	14	H42612
32	10.6	58.9	58	17	BH256481
33	10.4	57.8	27	17	AQ025667
c 34	10.4	57.8	52	9	AA485733
c 35	10.2	56.7	33	12	BF026570
c 36	10.2	56.7	34	9	A1192963
37	10.2	56.7	35	17	TA177A10Q
38	10.2	56.7	36	17	BH751662
39	10.2	56.7	36	17	BH751810
40	10.2	56.7	38	17	BH611852
41	10.2	56.7	38	17	BH618406
42	10.2	56.7	38	17	BH618437
43	10.2	56.7	38	17	BH618558
44	10.2	56.7	38	17	BH751595
45	10.2	56.7	39	13	BI838507

ALIGNMENTS

RESULT 1
TA58C07Q
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

TA58C07Q 49 bp DNA linear GSS 13-DEC-2000
T. brucei sheared genomic DNA clone 58c07, reverse sequence,
genomic survey sequence.
AL455710 GI:11857988
GSS.
Trypanosoma brucei.
Trypanosoma brucei
Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
1 (bases 1 to 49)
Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
Melville, S.E., Rajandream, M.A. and Barrell, B.G.
Direct Submission
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
Cambridge CB10 1SA, E-mail: barrellesanger.ac.uk and
nhlesanger.ac.uk
Constructed at the Institute for Genomic Research (TIGR),
Rockville, MD. Genomic DNA isolated from a cloned population of
Trypanosoma brucei (TREU927/4 GUFat 10.1) was mechanically sheared
to give a tight size distribution (4 kb). The v + i method used for the library construction is
described in detail in Smith, H. and Venter, J.C. (Making small
insert libraries for whole genome shotgun sequencing projects. In
Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.

Barrell, Oxford University Press, 1999)

Email: nelsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available at http://www.sanger.ac.uk/Projects/T_brucei/.

FEATURES

```
1. .49
/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="58c07"
```

SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 50)
 AUTHORS Suzuki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano
 TITLE Construction and characterization of a full length-enriched and
 JOURNAL a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
 COMMENT BodyMap: human gene expression database
 Unpublished (1995)
 Contact: Okubo, K.
 Institute for Molecular and Cellular Biol
 Osaka University
 1-3, Yamada-oka, Suita, Osaka Pref. 565, Japan
 Tel: 06-877-5111(ex.3315)
 Email: kousaku@imcb.osaka-u.ac.jp
 Human Gene Signature, 3'-directed cDNA sequence. We are not
 submitting the same cDNA sequence redundantly to DBJ since 1993.
 For the abundance information of clones with this sequence in this
 library and as well as in other 3'-directed libraries, see
 http://www.imcb.osaka-u.ac.jp/bodymap/. The sequences of the clones
 represented by this GS sequences is also found there.

FEATURES
 source
 1..50
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="Human adult (K.Okubo)"
 /dev_stage="adult"
 /note="Organ: blood; Vector: l-gt-11; Site: Eco-RI;
 Monocytos were prepared from blood by ficoll-hypaque,
 percoll and T cell rosetting purification steps (purity:
 96 %). mRNA was prepared from activated monocytes from a
 patient with rheumatoid arthritis. mRNA was reverse
 transcribed with MuLV. Using Eco-RI linkers cDNA was
 cloned into l-gt-11 vector arms. The cDNA library was
 screened by differential hybridization using radioactively
 marked ss-cDNA from activated and non-activated
 monocytes."

BASE COUNT 11 a 11 c 12 g 15 t 1 others
 ORIGIN
 Query Match 67.8%; Score 12.2; DB 14; Length 50;
 Best Local Similarity 64.7%; Pred. No. 4.6e+04;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AUGGCCUACGUGGCG 17
 ||:||||: ||: |||
 Db 34 AATGGCCCTCGATGCG 18

RESULT 5
 AUI05634 50 bp mRNA linear EST 30-AUG-2001
 LOCUS Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
 DEFINITION KAT08740, mRNA sequence.
 ACCESSION AUI05634
 VERSION AUI05634.1 GI:13555155
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 50)
 AUTHORS Suzuki, Y., Taira, H., Tsunoda, T., Mizushima-Sugano, J., Sese, J., Hata
 TITLE Diverse transcriptional initiation revealed by fine, large-scale
 JOURNAL mapping of mRNA start sites
 MEDLINE EMBO Rep. 2 (5), 388-393 (2001)
 COMMENT Contact: Yutaka Suzuki
 Department of Virology
 Institute of Medical Science, University of Tokyo
 4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
 Email: yusuzuki@ims.u-tokyo.ac.jp

Suzuki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano
 S. Construction and characterization of a full length-enriched and
 a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
 Location/Qualifiers
 1..50
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="KAT08740"
 /clone_lib="Sugano Homo sapiens cDNA library"
 /note="Differential display comparison of untreated and
 dimethylfumarate treated U937 cells"
 BASE COUNT 10 a 14 c 14 g 12 t
 ORIGIN
 Query Match 65.6%; Score 11.8; DB 9; Length 50;
 Best Local Similarity 66.7%; Pred. No. 7.2e+04;
 Matches 10; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 4 GGCCUACGUGGCGA 18
 |||:|:|:|:|:|:|
 Db 20 GCGTATCCGTGCGA 34

RESULT 6
 AV847604/c 56 bp mRNA linear EST 08-NOV-2001
 LOCUS AV847604 Nori Satoh unpublished cDNA library, larva Ciona
 DEFINITION intestinalis cDNA clone rcilv08b14 3', mRNA sequence.
 ACCESSION AV847604
 VERSION AV847604.1 GI:16828139
 KEYWORDS EST.
 SOURCE Ciona intestinalis.
 ORGANISM Ciona intestinalis
 Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
 Phlebobranchia; Clonidae; Ciona.
 REFERENCE 1 (bases 1 to 56)
 AUTHORS Satoh, N., Satou, Y., Kohara, Y. and Shin-i, T.
 TITLE Expressed genes in Ciona intestinalis
 JOURNAL Unpublished (2000)
 COMMENT Contact: Nori Satoh
 Department of Zoology
 Kyoto University
 Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
 Tel: 81-75-753-4081
 Fax: 81-75-705-1113
 Email: satoheascidian.zool.kyoto-u.ac.jp.
 Location/Qualifiers
 1..56
 /organism="Ciona intestinalis"
 /db_xref="taxon:7719"
 /clone_lib="rcilv08b14"
 /clone_lib="Nori Satoh unpublished cDNA library, larva"
 /tissue_type="whole animal"
 /dev_stage="larva"
 /note="vector: pBluescript SK"
 BASE COUNT 12 a 11 c 18 g 15 t
 ORIGIN
 Query Match 65.6%; Score 11.8; DB 10; Length 56;
 Best Local Similarity 73.3%; Pred. No. 7.4e+04;
 Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGCCUACGUGGCG 16
 |:|:|:|:|:|:|:|
 Db 45 ATGCCCAACGGTGC 31

RESULT 7
 AZ800642 29 bp DNA linear GSS 16-FEB-2001
 LOCUS AZ800642
 DEFINITION 2M0058G16R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC2M0058G16 R, DNA sequence.
 ACCESSION AZ800642

```

VERSION      AZ800642.1  GI:12952965
KEYWORDS     GSS.
SOURCE       Mus musculus
ORGANISM     house mouse.

REFERENCE    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
1            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
AUTHORS     Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
            Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
            M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
            and Wright,D., Weiss,R.

TITLE       Mouse whole genome scaffolding with paired end reads from 10kb
            plasmid inserts

JOURNAL      Unpublished (2000)
COMMENT      Contact: Robert B. Weiss
            University of Utah Genome Center
            Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
            84112, USA
            Tel: 801 585 5606
            Fax: 801 585 7177
            Email: ddunn@genetics.utah.edu
            Insert Length: 10000 Std Error: 0.00
            Plate: 0058 row: G column: 16
            Seq primer: CACACAGGAAACAGCTATGACC
            Class: plasmid ends
            High quality sequence stop: 29.

FEATURES     Location/Qualifiers
             1..29
                /organism="Mus musculus"
                /strain="C57BL/6J"
                /db_xref="taxon:10090"
                /clone="UUGC2M0058G16"
                /clone_lib="Mouse 10kb plasmid UUGCLM library"
                /sex="Male"
                /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
                /note="Vector: PWD42nv; Purified genomic DNA from M.
            musculus C57BL/6J (male) was obtained from the Jackson
            Laboratory Mouse DNA Resource
            (http://www.jax.org/resources/documents/dnares/). The DNA
            was hydrodynamically sheared by repeated passage through a
            0.005 inch orifice at constant velocity. The sheared DNA
            was blunt end-repaired with T4 DNA polymerase and T4
            polynucleotide kinase. Adaptor oligonucleotides were
            ligated to the blunt ends in high molar excess. The
            adapted DNA was purified and size-selected for a 9.5 to
            10.5 kb range using preparative agarose gel
            electrophoresis. Vector DNA was prepared from a derivative
            of PWD42 (gi14732114|gb|AF129072.1), a copy-number
            inducible derivative of plasmid p1. The vector was ligated
            with adaptors complementary to the insert adaptors and
            purified. The sheared, adapted mouse DNA was annealed to
            adapted vector DNA, and transformed into
            chemically-competent E. coli XL10-Gold (Stratagene) cells
            and selected for ampicillin resistance."
             4 a 5 c 7 g 13 t

BASE COUNT   4 a 5 c 7 g 13 t
ORIGIN

Query Match      64.4%; Score 11.6; DB 17; Length 29;
Best Local Similarity 61.1%; Pred. No. 7.8e+04;
Matches 11; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUACUGGUGCGCA 18
    |||||:|:|:|:|
Db 2 ATGGCCTATGGGCGCA 19

RESULT 8
AA863171
LOCUS      55 bp mRNA linear EST 29-APR-1998
DEFINITION o91e02.s1 NCI-CGAP_Ki5 Homo sapiens cDNA clone IMAGE:1455674 3'
            similar to SW:H1E3_HUMAN Q15738 H105E3 PROTEIN. ; mRNA sequence.
ACCESSION  AA863171

```

```

VERSION      AA863171.1  GI:2955650
KEYWORDS     EST.
SOURCE       human.
ORGANISM     Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE    1 (bases 1 to 55)
AUTHORS     NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
            Unpublished (1997)
COMMENT      Contact: Robert Strausberg, Ph.D.
            Email: cgaps-r@mail.nih.gov
            Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
            Emert-Buck, M.D., Ph.D.
            CDNA Library Preparation: M. Bento Soares, Ph.D.
            CDNA Library Arrayed by: Greg Lennon, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
Insert Length: 360 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 1.

FEATURES     Location/Qualifiers
             1..55
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="IMAGE:1455674"
                /clone_lib="NCI-CGAP_Ki5"
                /tissue_type="2 pooled tumors (clear cell type)"
                /lab_host="DH10B"
                /note="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with
            a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
            strand cDNA was primed with a Not I - oligo(dT) primer [5'
            AACTGGAGAAATTCGGCGCGCAATATATTTTTTTTTTTT 3'],
            double-stranded cDNA was ligated to Eco RI adaptors
            (Pharmacia), digested with Not I and cloned into the Not I
            and Eco RI sites of the modified pT7T3 vector. Library
            went through one round of normalization. Library
            constructed by Bento Soares and M. Fatima Bonaldo. "
             12 a 16 c 15 g 12 t

BASE COUNT   12 a 16 c 15 g 12 t
ORIGIN

Query Match      64.4%; Score 11.6; DB 9; Length 55;
Best Local Similarity 66.7%; Pred. No. 9.2e+04;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUACUGGUGCGCA 18
    ||:|||||:|:|:|:|
Db 28 AATTGCCAACACGATGCGA 45

RESULT 9
AZ840293/c
LOCUS      30 bp DNA linear GSS 20-FEB-2001
DEFINITION 2M0136H17R Mouse 10kb plasmid UUGCLM library Mus musculus genomic
            clone UUGC2M0136H17 R, DNA sequence.
ACCESSION  AZ840293
VERSION     AZ840293.1  GI:13010201
KEYWORDS    GSS.
SOURCE      house mouse.
ORGANISM    Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
            1 (bases 1 to 30)
            Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
            Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
            M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
            and Wright,D., Weiss,R.

REFERENCE    1
AUTHORS     Mouse whole genome scaffolding with paired end reads from 10kb

```


JOURNAL
COMMENT

plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0136 row: H column: 17
Seq primer: CACACGGAACACCTATGACC
Class: plasmid ends
High quality sequence stop: 30.

FEATURES
source

1. .30
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUC2M0136H17"
/clone_lib="Mouse 10kb plasmid UUC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57Bl/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 (gi14732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN

4 a 10 c 9 g 7 t
Query Match 62.2%; Score 11.2; DB 17; Length 30;
Best Local Similarity 62.5%; Pred. No. 1.2e+05;
Matches 10; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
Qy 1 AAUGGCCUUAUGCGUGC 16
Db 30 AATGGCCTGTCCGAGC 15

RESULT 10
AZ605771/c

LOCUS
DEFINITION
36 bp DNA linear GSS 13-DEC-2000
clone UUC1M0427K13 F, DNA sequence.
ACCESSION
AZ605771
VERSION
AZ605771.1 GI:11727961
KEYWORDS
GSS.
SOURCE
Mus musculus
house mouse.
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 36)
Dunn, B., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A., and Wright, D., Weiss, R.
TITLE
Mouse whole genome scaffolding with paired end reads from 10kb

JOURNAL
COMMENT

plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0427 row: K column: 13
Seq primer: CGTTGTAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 36.

FEATURES
source

1. .36
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUC1M0427K13"
/clone_lib="Mouse 10kb plasmid UUC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57Bl/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 (gi14732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN

14 a .10 c 5 g 7 t
Query Match 62.2%; Score 11.2; DB 17; Length 36;
Best Local Similarity 56.2%; Pred. No. 1.3e+05;
Matches 9; Conservative 4; Mismatches 3; Indels 0; Gaps 0;
Qy 2 AUGGCCUUAUGCGUGCG 17
Db 18 ATGGTCTATCAGTGTG 3

RESULT 11
AA931624

LOCUS
DEFINITION
37 bp mRNA linear EST 24-APR-1998
oo35b08.s1 NCI CGAP Lu5 Homo sapiens cDNA clone IMAGE:1568151 3', similar to SW:IBA2_HUMAN Q14657 ITBA2 PROTEIN ;, mRNA sequence.
ACCESSION
AA931624
VERSION
AA931624.1 GI:3086010
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 37)
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE
National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL
Unpublished (1997)
CONTACT: Robert Strausberg, Ph.D.

Email: ccapbs-remail.nth.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Suck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CCAP clone distribution
Information can be found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 1.

FEATURES	SOURCE
1. Feature 1	Source 1
2. Feature 2	Source 2
3. Feature 3	Source 3
4. Feature 4	Source 4
5. Feature 5	Source 5
6. Feature 6	Source 6
7. Feature 7	Source 7
8. Feature 8	Source 8
9. Feature 9	Source 9
10. Feature 10	Source 10
11. Feature 11	Source 11
12. Feature 12	Source 12
13. Feature 13	Source 13
14. Feature 14	Source 14
15. Feature 15	Source 15
16. Feature 16	Source 16
17. Feature 17	Source 17
18. Feature 18	Source 18
19. Feature 19	Source 19
20. Feature 20	Source 20
21. Feature 21	Source 21
22. Feature 22	Source 22
23. Feature 23	Source 23
24. Feature 24	Source 24
25. Feature 25	Source 25
26. Feature 26	Source 26
27. Feature 27	Source 27
28. Feature 28	Source 28
29. Feature 29	Source 29
30. Feature 30	Source 30
31. Feature 31	Source 31
32. Feature 32	Source 32
33. Feature 33	Source 33
34. Feature 34	Source 34
35. Feature 35	Source 35
36. Feature 36	Source 36
37. Feature 37	Source 37
38. Feature 38	Source 38
39. Feature 39	Source 39
40. Feature 40	Source 40
41. Feature 41	Source 41
42. Feature 42	Source 42
43. Feature 43	Source 43
44. Feature 44	Source 44
45. Feature 45	Source 45
46. Feature 46	Source 46
47. Feature 47	Source 47
48. Feature 48	Source 48
49. Feature 49	Source 49
50. Feature 50	Source 50
51. Feature 51	Source 51
52. Feature 52	Source 52
53. Feature 53	Source 53
54. Feature 54	Source 54
55. Feature 55	Source 55
56. Feature 56	Source 56
57. Feature 57	Source 57
58. Feature 58	Source 58
59. Feature 59	Source 59
60. Feature 60	Source 60
61. Feature 61	Source 61
62. Feature 62	Source 62
63. Feature 63	Source 63
64. Feature 64	Source 64
65. Feature 65	Source 65
66. Feature 66	Source 66
67. Feature 67	Source 67
68. Feature 68	Source 68
69. Feature 69	Source 69
70. Feature 70	Source 70
71. Feature 71	Source 71
72. Feature 72	Source 72
73. Feature 73	Source 73
74. Feature 74	Source 74
75. Feature 75	Source 75
76. Feature 76	Source 76
77. Feature 77	Source 77
78. Feature 78	Source 78
79. Feature 79	Source 79
80. Feature 80	Source 80
81. Feature 81	Source 81
82. Feature 82	Source 82
83. Feature 83	Source 83
84. Feature 84	Source 84
85. Feature 85	Source 85
86. Feature 86	Source 86
87. Feature 87	Source 87
88. Feature 88	Source 88
89. Feature 89	Source 89
90. Feature 90	Source 90
91. Feature 91	Source 91
92. Feature 92	Source 92
93. Feature 93	Source 93
94. Feature 94	Source 94
95. Feature 95	Source 95
96. Feature 96	Source 96
97. Feature 97	Source 97
98. Feature 98	Source 98
99. Feature 99	Source 99
100. Feature 100	Source 100

1. 37
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1568151"
/clone_lib="NCI_CGAP_Lu5"
/tissue_type="cardinoid"
/lab_host="DH108"
/notes="Organ: lung; Vector: pT7T3p-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from neuroendocrine lung carcinoid, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library is normalized. Library was constructed by Bento Soares and M. Fatima Bonaldo."
10 c 11 q 9 t

BASE COUNT
ORIGIN

Query Match 62.2%; Score 11.2; DB 9; Length 37;

RESULT 12					
AZ317769/c					
LOCUS					
DEFINITION	AZ317769	45 bp	DNA	linear	GSS 29-SEP-2000
	1M0036002R	Mouse 10kb	plasmid	UUGC1M library	Mus musculus genomic
		clone	UUGC1M0036002 R,	DNA sequence.	

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 45)

REFERENCE
AUTHORS

Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duvall, B., Hamil, C.,
Islam, H., Longacre, S., Mahmood, M., Meenen, E., Pedersen, T., Rellly
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.

TITLE	Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts and Wright, D. B. Weiss, R. A.
JOURNAL	Unpublished (2000)
COMMENT	Contact: Robert B. Weiss University of Utah Genome Center University of Utah Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA Tel: 801 585 5606 Fax: 801 585 7177 Email: ddunn@genetics.utah.edu Insert Length: 10000 Std Error: 0.00 Plate: 0036 row: 0 column: 02 Seq primer: CACACAGGAACAGCTATGACC Class: plasmid ends

High quality sequence stop: 45.
Location/Qualifiers
1. .45

```

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0036002"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="F. Coli strain XL10-Gold, T1-resistant, F-"
/notes="Vector: pWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (g1147321141gb1AF129072.1), a copy-number inducible derivative of plasmid p1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

```

BASE COUNT	10 a	11 c	12 g	12 t	
ORIGIN					
Query Match		62.2%	Score 11.2;	DB 17;	Length 45;
Best Local Similarity		56.2%;	Pred. No. 1.4e+05;		
Matches	9;	Conservative	4;	Mismatches	3; Indels 0; Gaps 0;
QY	1	AAUAGCCCUAUCGGUGC	16		
		11:1	1:1:1:1	:	1:1
Db	37	AATGCTCATCGATGC	22		

RESULT 13					
BH618002					
LOCUS	BH618002	51 bp	DNA	linear	GSS 30-JAN-2002
DEFINITION	SALK_038348 Arabidopsis thaliana TDNA insertion lines Arabidopsis thaliana genomic clone SALK_038348, DNA sequence.				

COORDINATOR
ORGANISM
Arabiidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabioidopsis.
1 (bases 1 to 51)
Alonso, J. M., Leisse, T. J., Barajas, P., Chen, H., Cheuk, R., Gadrinab,
C., Jeske, A., Karnes, M., Kim, C. J., Parker, H., Prednits, L., Shinn, P.,
Zimmerman, J. and Ecker, J. R.

FEATURES
source

/organism="Arabidopsis thaliana"
 /strain="Columbia 0"
 /db_xref="taxon:3702"
 /clone="SALK_038348"
 /clone_lib="Arabidopsis thaliana TDNA insertion lines"

/note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT 10 a 15 c 17 g 9 t

ORIGIN

Query Match 62.2%; Score 11.2; DB 17; Length 51;

Best Local Similarity 56.2%; Pred. No. 1.4e+05; Mismatches 9; Conservative 4; Indels 0; Gaps 0;

Qy 3 UGCCCUAUCGUGCGA 18

Db 29 TGCCCTATAGTGGGA 44

RESULT 14

AA721034

LOCUS

DEFINITION nx89h04.sl NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1269463 3' similar to WP:F56DI.3 CE01971 ;, mRNA sequence.

ACCESSION AA721034

VERSION AA721034.1

KEYWORDS EST

SOURCE human.

ORGANISM Homo sapiens

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

52 bp mRNA linear EST 22-JAN-1998
 NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 Email: cgaps-r@mail.nih.gov
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
 Ph.D., Gerald Marti, M.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
 Bonaldo, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 730 Std Error: 0.00

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 1.

Location/Qualifiers

1..52

FEATURES

source

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1269463"

/clone_lib="NCI_CGAP_GCB1"

/tissue_type="germinal center B cell"

/lab_host="DH10B"

/note="vector: pT73b-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was prepared from human tonsillar cells enriched for germinal center B cells by flow sorting (CD20+, Igu-), provided by Dr. Louis M. Staudt (NCI), Dr. David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was primed with a Not I - oligo(dT) primer [5'-TGTTACCACTCTCAAGTGGGAGCGGCTCATTTTTTTTTTTT-3']. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library

BASE COUNT 12 a 12 c 21 g 7 t

ORIGIN

Query Match

Best Local Similarity 62.2%; Score 11.2; DB 9; Length 52;

Mismatches 11; Conservative 2; Indels 0; Gaps 0;

Qy 1 A AUGGCCUACGUGGC 16

Db 12 AACGCCGATGTGGTC 27

RESULT 15

BH641243

LOCUS

DEFINITION BH641243 1008046D05.2EL_y1 1008 - RescueMu Grid I Zea mays genomic, DNA sequence.

ACCESSION BH641243

VERSION BH641243.1

KEYWORDS GSS.

SOURCE ze mays.

ORGANISM ze mays

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Maize genomic sequences found using engineered RescueMu transposon
 Unpublished (2001)
 Contact: Walbot V

Department of Biological Sciences

Stanford University

855 California Ave, Palo Alto, CA 94304, USA

Tel: 650 723 2227

Fax: 650 725 8221

Email: walbot@stanford.edu

Possible ligation site of ends cut by 2 different endonucleases.
 Reverse complemented post-ligation sequence from source sequence.
 Plate: 1008046 row: 17

Class: transposon-tagged.

Location/Qualifiers

1..54

FEATURES

source

/organism="Zea mays"

/cultivar="mixed background W23/A188/B73"

/db_xref="taxon:4577"

/clone_lib="1008 - RescueMu Grid I"

/tissue_type="leaf"

/dev_stage="adult"

/lab_host="DH10B"

/note="Organ: leaf; Vector: RescueMu (engineered from

pBlueScript backbone); Site_1: BamHI; Site_2: BglII;

RescueMu is a 4.9 kb, modified maize Mu transposon

designed to allow plasmid rescue from total genomic DNA.

Mu elements insert preferentially into transcription

units. For more information on RescueMu, go to the web

site www.zmmb.iastate.edu and follow the links for

'RescueMu.' Grid I was grown at Berkeley in 2001. DNA was

extracted from leaf punches, double digested using BamHI

and BglII, and ligated to form circular plasmids. DH10B

cells were transformed and then screened on LB plates with

ampicillin."

BASE COUNT 12 a 12 c 17 g 13 t

ORIGIN

Query Match

Best Local Similarity 62.2%; Score 11.2; DB 17; Length 54;

Mismatches 11; Conservative 2; Indels 0; Gaps 0;

Qy 2 AUGGCCUACGUGGC 17

Db 31 ATGGCCTTACGGGCG 46

Search completed: July 6, 2003, 15:28:29
Job time : 1212.27 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:33:06 ; Search time 47.4545 seconds
(without alignments)
116.326 Million cell updates/sec

Title: US-09-780-929-98

Perfect score: 18

Sequence: 1 aauggccuacgugcgca 18

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 635134

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents NA: *
1: /cgn2_6/ptodata/1/ina/5A_COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	12.2	67.8	29	4	US-09-199-637A-144
2	11.8	65.6	30	1	US-07-997-133-5
3	11.8	65.6	30	5	US-07-997-133-5
4	11.8	65.6	40	4	US-09-626-929-3
5	11.8	65.6	40	4	US-09-484-850-3
6	11.8	65.6	40	4	US-09-408-392-3
7	11.8	65.6	40	4	US-09-626-930-3
8	11.8	65.6	40	4	US-09-626-528-3
9	11.6	64.4	26	1	US-08-054-480-5
10	11.6	64.4	29	1	US-08-413-803-10
11	11.6	64.4	29	1	US-08-321-488A-10
12	11.6	64.4	29	5	PCT-US95-03776-10
13	11.6	64.4	30	1	US-08-413-803-9
14	11.6	64.4	30	1	US-08-321-488A-9
15	11.6	64.4	30	2	US-08-943-915-14
16	11.6	64.4	30	4	US-08-482-918-36
17	11.6	64.4	30	4	US-09-224-681-36
18	11.6	64.4	30	4	US-08-336-728A-36
19	11.6	64.4	30	5	PCT-US95-03776-9
20	11.6	64.4	41	2	US-08-781-620B-10
21	11.6	64.4	46	4	US-08-520-678A-15
22	11.6	64.4	46	4	US-08-897-126-15
23	11.4	63.3	17	2	US-08-292-620A-1886
24	11.4	63.3	17	2	US-08-292-620A-1886
25	11.4	63.3	17	3	US-09-071-845-1886
26	11.4	63.3	17	3	US-09-071-845-1886
27	11.4	63.3	21	4	US-08-397-220B-24

Sequence 24, Appl
Sequence 24, Appl
Sequence 35, Appl
Sequence 35, Appl
Sequence 35, Appl
Sequence 35, Appl
Sequence 35, Appl
Sequence 1095, Ap
Sequence 7, Appl
Sequence 31, Appl
Sequence 19, Appl
Sequence 26, Appl
Sequence 11, Appl
Sequence 11, Appl
Sequence 1, Appl
Sequence 75, Appl
Sequence 75, Appl
Sequence 1, Appl

28 11.4 63.3 21 4 US-08-650-093C-24
29 11.4 63.3 21 4 US-08-823-895A-24
30 11.4 63.3 25 3 US-09-150-133-35
31 11.4 63.3 25 3 US-09-150-141-35
32 11.4 63.3 25 4 US-09-374-493-35
33 11.4 63.3 25 4 US-09-374-824-35
34 11.4 63.3 25 4 US-09-374-492-35
35 11.4 63.3 47 4 US-09-641-638-1095
36 11.2 62.2 24 5 PCT-US96-09473-7
37 11.2 62.2 26 2 US-08-805-918-31
38 11.2 62.2 26 2 US-08-811-028-19
39 11.2 62.2 30 1 US-08-393-985-26
40 11.2 62.2 30 4 US-09-105-697-11
41 11.2 62.2 30 4 US-09-146-631-11
42 11.2 62.2 33 5 PCT-US96-09473-1
43 11.2 62.2 40 2 US-08-425-684-75
44 11.2 62.2 40 2 US-08-675-502-75
45 11.2 62.2 43 4 US-09-686-179A-1

ALIGNMENTS

RESULT 1

US-09-199-637A-144/c
; Sequence 144, Application US/09199637A

; Patent No. 635411

; GENERAL INFORMATION:

; APPLICANT: Ausubel, Frederick

; APPLICANT: Goodman, Howard M.

; APPLICANT: Rahme, Laurence G.

; APPLICANT: Mahajan-Miklos, Shalina

; APPLICANT: Tan, Man-Wah

; APPLICANT: Cao, Hui

; APPLICANT: Drenkard, Eliana

; APPLICANT: Tsongalis, John

; TITLE OF INVENTION: VIRULENCE-ASSOCIATED NUCLEIC ACID

; TITLE OF INVENTION: SEQUENCES AND USES THEREOF

; FILE REFERENCE: 00786/361002

; CURRENT APPLICATION NUMBER: US/09/199,637A

; CURRENT FILING DATE: 1998-11-25

; PRIOR APPLICATION NUMBER: 60/066,517

; PRIOR FILING DATE: 1997-11-25

; NUMBER OF SEQ ID NOS: 437

; SOFTWARE: FastSeq for Windows Version 4.0

; SEQ ID NO 144

; LENGTH: 29

; TYPE: DNA

; ORGANISM: Pseudomonas aeruginosa

US-09-199-637A-144

Query Match 67.8%; Score 12.2; DB 4; Length 29;

Best Local Similarity 64.7%; Pred. No. 2.6e+02;

Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAUGGCCUACGUGCG 17

11 111 :1:11 :111

Db 19 AACGGCGTATCGTTGCG 3

RESULT 2

US-07-997-133-5

; Sequence 5, Application US/07997133

; Patent No. 5288855

; GENERAL INFORMATION:

; APPLICANT: Bergonzoni, Laura

; APPLICANT: Mazue, Guy

; APPLICANT: Isacchi, Antonella

; APPLICANT: Roncucci, Romeo

; APPLICANT: Sarmientos, Paolo

; TITLE OF INVENTION: Extracellular Form of the Human

; TITLE OF INVENTION: Fibroblast Growth Factor Receptor

; NUMBER OF SEQUENCES: 8

```
;
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 Jefferson Davis Highway, Fourth Floor
; CITY: Arlington
; STATE: Virginia
; ZIP: 22202
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/997,133
; FILING DATE: 28-DEC-1992
; CLASSIFICATION: 530
;
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: US/07/642,755
; FILING DATE: 18-JAN-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Oblon, No. 528885man F.
; REGISTRATION NUMBER: 24,618
; REFERENCE/DOCKET NUMBER: 769-226-0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703)521-4500
; TELEFAX: (703)486-2347
; TELEX: 248855 OPAT UR
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-07-997-133-5

Query Match 65.6%; Score 11.8; DB 1; Length 30;
Best Local Similarity 73.3%; Pred. No. 4.4e+02;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUGCGGUC 16
Db 1 ACGGCCTAGCGGTGC 15

RESULT 3
US-07-997-133-5
; Sequence 5, Application US/07997133
; GENERAL INFORMATION:
; APPLICANT: Bergonzoni, Laura
; APPLICANT: Mazue, Guy
; APPLICANT: Isacchi, Antonella
; APPLICANT: Roncucci, Romeo
; APPLICANT: Samientos, Paolo
; TITLE OF INVENTION: Extracellular Form of the Human
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 Jefferson Davis Highway, Fourth Floor
; CITY: Arlington
; STATE: Virginia
; ZIP: 22202
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/997,133
; FILING DATE: 28-DEC-1992
; CLASSIFICATION: 530
```

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;
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/07/642,755
; FILING DATE: 18-JAN-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Oblon, Norman F.
; REGISTRATION NUMBER: 24,618
; REFERENCE/DOCKET NUMBER: 769-226-0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703)521-4500
; TELEFAX: (703)486-2347
; TELEX: 248855 OPAT UR
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-07-997-133-5

Query Match 65.6%; Score 11.8; DB 5; Length 30;
Best Local Similarity 73.3%; Pred. No. 4.4e+02;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUGCGGUC 16
Db 1 ACGGCCTAGCGGTGC 15

RESULT 4
US-09-626-929-3
; Sequence 3, Application US/09626929
; Patent No. 6319714
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILHELM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,929
; 2000-07-27
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
; US-09-626-929-3

Query Match 65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUGCGGUC 15
Db 6 ACTGGCGTATCGGTG 20
```

```
RESULT 5
US-09-484-850-3
; Sequence 3, Application US/09484850
; Patent No. 6368861
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/484,850
; CURRENT FILING DATE: 2000-01-18
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-484-850-3
Query Match 65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
   | :||| :|:|:|
Db 6 ACTGGCGTATCGGTG 20

RESULT 6
US-09-408-392-3
; Sequence 3, Application US/09408392
; Patent No. 6376246
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/408,392
; CURRENT FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-408-392-3
Query Match 65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
   | :||| :|:|:|
Db 6 ACTGGCGTATCGGTG 20

RESULT 7
US-09-626-930-3
; Sequence 3, Application US/09626930
; Patent No. 6423542
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,930
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-626-930-3
Query Match 65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
   | :||| :|:|:|
Db 6 ACTGGCGTATCGGTG 20

RESULT 8
US-09-626-528-3
; Sequence 3, Application US/09626528
; Patent No. 6426224
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,528
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
```

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US-09-408-392-3
Query Match 65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
   | :||| :|:|:|
Db 6 ACTGGCGTATCGGTG 20

RESULT 7
US-09-626-930-3
; Sequence 3, Application US/09626930
; Patent No. 6423542
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,930
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-626-930-3
Query Match 65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
   | :||| :|:~|:|
Db 6 ACTGGCGTATCGGTG 20

RESULT 8
US-09-626-528-3
; Sequence 3, Application US/09626528
; Patent No. 6426224
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,528
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
```

;; PRIOR APPLICATION NUMBER: 60/118,813
;; PRIOR FILING DATE: 1999-02-05
;; PRIOR APPLICATION NUMBER: 60/141,049
;; PRIOR FILING DATE: 1999-06-24
;; NUMBER OF SEQ ID NOS: 26
;; SOFTWARE: PatentIn ver. 2.1
;; SEQ ID NO 3
;; LENGTH: 40
;; TYPE: DNA
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence: Bridging
;; OTHER INFORMATION: oligonucleotides
US-09-626-528-3

Query Match 65.68; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.08; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
| :| | | :| :| | | :|
Db 6 ACTGGCGTATCGGTG 20

RESULT 9
US-08-054-480-5
;; Sequence 5, Application US/08054480
;; Patent No. 5525504
;; GENERAL INFORMATION:
;; APPLICANT: Goebel, Werner
;; APPLICANT: Libby, Stephen
;; APPLICANT: Heffron, Fred
;; TITLE OF INVENTION: CYTOLYSIN GENE AND GENE PRODUCT
;; NUMBER OF SEQUENCES: 5
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: MILLEN, WHITE, ZELANO, & BRANIGAN, P.C.
;; STREET: 2200 CLARENDON BOULEVARD, SUITE 1400
;; CITY: ARLINGTON
;; STATE: VIRGINIA
;; COUNTRY: USA
;; ZIP: 22201

;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.25
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/054,480
;; FILING DATE: 04-APR-1993
;; CLASSIFICATION: 435
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Heaney, Brian P.
;; REGISTRATION NUMBER: 32,542
;; REFERENCE/DOCKET NUMBER: MERCK 1496
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 703 243 6333
;; TELEFAX: 703 243 6410
;; TELEX: 64191

;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 26 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; ORIGINAL SOURCE:
;; ORGANISM: SALMONELLA
US-08-054-480-5

Query Match 64.4%; Score 11.6; DB 1; Length 26;
Best Local Similarity 66.7%; Pred. No. 5.7e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
| :| | | | | :| :| | | | | :| :| | | | |
Db 7 AATGGCAGAGAGGTGCGA 24

RESULT 10

US-08-413-803-10/c
;; Sequence 10, Application US/08413803
;; Patent No. 5766581
;; GENERAL INFORMATION:
;; APPLICANT: Bartley, Timothy D.
;; APPLICANT: Bogenberger, Jakob M.
;; APPLICANT: Bosselman, Robert A.
;; APPLICANT: Hunt, Pamela
;; APPLICANT: Kinstler, Olaf B.
;; APPLICANT: Samal, Babru B.
;; TITLE OF INVENTION: METHODS FOR TREATING MAMMALS WITH
;; TITLE OF INVENTION: MONO-PEGYLATED PROTEINS THAT STIMULATE MEGAKARYOCYTE
;; TITLE OF INVENTION: GROWTH AND DIFFERENTIATION
;; NUMBER OF SEQUENCES: 34
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: AMGEN INC.
;; STREET: 1840 DeHavilland Drive
;; CITY: Thousand Oaks
;; STATE: California
;; COUNTRY: US
;; ZIP: 91320-1789

;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/413,803
;; FILING DATE: 30-MAR-1995
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/221,768
;; FILING DATE: 31-MAR-1994
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/252,628
;; FILING DATE: 31-MAY-1994
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/321,488
;; FILING DATE: 12-OCT-1994
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/347,780
;; FILING DATE: 30-NOV-1994
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Cook Ph.D., Robert R.
;; REGISTRATION NUMBER: 31,602
;; REFERENCE/DOCKET NUMBER: A-290D
;; INFORMATION FOR SEQ ID NO: 10:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 29 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: cdna
US-08-413-803-10

Query Match 64.4%; Score 11.6; DB 1; Length 29;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
| :| | | | | :| :| | | | | :| :| | | | |
Db 19 AAAGGCGTATCCGGCGA 2

RESULT 11
US-08-321-488A-10/c
;; Sequence 10, Application US/08321488A


```
; Patent No. 5795569
; GENERAL INFORMATION:
; APPLICANT: Bartley, Timothy D.
; APPLICANT: Bogenberger, Jakob M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Hunt, Pamela
; APPLICANT: Kinstler, Olaf B.
; APPLICANT: Samal, Babru B.
; TITLE OF INVENTION: MONO-PEGYLATED PROTEINS THAT STIMULATE
; TITLE OF INVENTION: MEGAKARYOCYTE GROWTH AND DIFFERENTIATION
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: AMGEN INC.
; STREET: 1840 DeHavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: US
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/321,488A
; FILING DATE: 12-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/252,628
; FILING DATE: 31-MAY-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/221,768
; FILING DATE: 31-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Cook, Robert R.
; REGISTRATION NUMBER: 31,602
; REFERENCE/DOCKET NUMBER: A-290B
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 29 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; US-08-321-488A-10

Query Match 64.4%; Score 11.6; DB 1; Length 29;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
Db 19 AAAGGCGCTATCCGGCCGA 2

RESULT 12
PCT-US95-03776-10/c
; Sequence 10, Application PC/TUS9503776
; GENERAL INFORMATION:
; APPLICANT: AMGEN INC.
; TITLE OF INVENTION: Compositions and Methods for Stimulating
; TITLE OF INVENTION: Megakaryocyte Growth and Differentiation
; NUMBER OF SEQUENCES: 34
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Amgen Inc.
; STREET: 1840 Dehavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: USA
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
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; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/03776
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Cook, Robert R.
; REFERENCE/DOCKET NUMBER: A-290-C
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 29 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; PCT-US95-03776-10

Query Match 64.4%; Score 11.6; DB 5; Length 29;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
Db 19 AAAGGCGCTATCCGGCCGA 2

RESULT 13
US-08-413-803-9/c
; Sequence 9, Application US/08413803
; Patent No. 5766581
; GENERAL INFORMATION:
; APPLICANT: Bartley, Timothy D.
; APPLICANT: Bogenberger, Jakob M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Hunt, Pamela
; APPLICANT: Kinstler, Olaf B.
; APPLICANT: Samal, Babru B.
; TITLE OF INVENTION: METHODS FOR TREATING MAMMALS WITH
; TITLE OF INVENTION: MONO-PEGYLATED PROTEINS THAT STIMULATE MEGAKARYOCYTE
; NUMBER OF SEQUENCES: 34
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: AMGEN INC.
; STREET: 1840 DeHavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: US
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/413,803
; FILING DATE: 30-MAR-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/221,768
; FILING DATE: 31-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/252,628
; FILING DATE: 31-MAY-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/321,488
; FILING DATE: 12-OCT-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/347,780
; FILING DATE: 30-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Cook Ph.D., Robert R.
; REGISTRATION NUMBER: 31,602
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REFERENCE/DOCKET NUMBER: A-290D
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-413-803-9

Query Match 64.4%; Score 11.6; DB 1; Length 30;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGUGCGCA 18
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Db 19 AAAGGCCTATCCGGCGCA 2

RESULT 14
US-08-321-488A-9/c
Sequence 9, Application US/08321488A
Patent No. 5795569
GENERAL INFORMATION:
APPLICANT: Bartley, Timothy D.
APPLICANT: Bogenberger, Jakob M.
APPLICANT: Bosselman, Robert A.
APPLICANT: Hunt, Pamela
APPLICANT: Kinstler, Olaf B.
APPLICANT: Samal, Babru B.
TITLE OF INVENTION: MEGA-PGYLATED PROTEINS THAT STIMULATE
TITLE OF INVENTION: MEGAKARYOCYTE GROWTH AND DIFFERENTIATION
NUMBER OF SEQUENCES: 27
CORRESPONDENCE ADDRESS:
ADDRESSEE: AMGEN INC.
STREET: 1840 DeHavilland Drive
CITY: Thousand Oaks
STATE: California
COUNTRY: US
ZIP: 91320-1789
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/321.488A
FILING DATE: 12-OCT-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/252,628
FILING DATE: 31-MAY-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/221,768
FILING DATE: 31-MAR-1994
ATTORNEY/AGENT INFORMATION:
NAME: Cook, Robert R.
REGISTRATION NUMBER: 31,602
REFERENCE/DOCKET NUMBER: A-290B
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-321-488A-9

Query Match 64.4%; Score 11.6; DB 1; Length 30;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGUGCGCA 18

Db 19 AAAGGCCTATCCGGCGCA 2
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RESULT 15
US-08-943-915-14/c
Sequence 14, Application US/08943915
Patent No. 5998170
GENERAL INFORMATION:
APPLICANT: Itoh, No. 5998170uyuki
APPLICANT: Martin, Frank
APPLICANT: Danilenko, Dmitry
TITLE OF INVENTION: A FIBROBLAST GROWTH FACTOR
NUMBER OF SEQUENCES: 33
CORRESPONDENCE ADDRESS:
ADDRESSEE: Amgen Inc.
STREET: 1840 DeHavilland Drive
CITY: Thousand Oaks
STATE: California
COUNTRY: USA
ZIP: 91320-1789
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/943,915
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mazza, Richard J.
REGISTRATION NUMBER: 27,657
REFERENCE/DOCKET NUMBER: A-469
TELECOMMUNICATION INFORMATION:
TELEPHONE: 805.447.4112
TELEFAX: 805.447.1090
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "OLIGONUCLEOTIDE"

US-08-943-915-14
Query Match 64.4%; Score 11.6; DB 2; Length 30;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

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Db 19 AAAGGCCTATCCGGCGCA 2

Search completed: July 6, 2003, 15:30:04
Job time : 48.4545 secs

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OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 15:04:31 ; Search time 102 Seconds
(without alignments)
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Title: US-09-780-929-98
Perfect score: 18
Sequence: 1 aaugccuauccggucga 18

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Gapop 10.0 , Gapext 1.0

Searched: 1085931 seqs, 780495707 residues

Total number of hits satisfying chosen parameters: 805896

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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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- 13: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*
- 14: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	18	100.0	29	10	US-09-780-929-107
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4	13.8	76.7	30	10	US-09-798-675-8
5	12.2	67.8	29	9	US-09-975-719-144
6	11.8	65.6	24	9	US-09-940-185-3899
7	11.6	64.4	30	10	US-09-005-243-36
8	11.6	64.4	30	10	US-09-224-683-36
9	11.6	64.4	46	9	US-09-880-508-15
10	11.6	64.4	46	9	US-10-158-314-15
11	11.6	64.4	50	9	US-09-741-179A-10
12	11.6	64.4	50	9	US-09-741-179A-14
13	11.4	63.3	24	9	US-09-940-185-3596
14	11.4	63.3	41	10	US-09-759-272B-4
15	11.4	63.3	42	10	US-09-838-386-14
16	11.4	63.3	45	10	US-09-838-386-13
17	11.4	63.3	45	10	US-09-838-386-17
18	11.4	63.3	45	10	US-09-838-386-18
19	11.4	63.3	51	9	US-10-211-088-46

c	20	11.2	62.2	17	9	US-09-780-533A-633	Sequence 633, Appl
c	21	11.2	62.2	19	9	US-09-796-081-3	Sequence 3, Appli
c	22	11.2	62.2	19	9	US-09-796-081-4	Sequence 4, Appli
c	23	11.2	62.2	22	10	US-09-997-664-85	Sequence 85, Appl
c	24	11.2	62.2	25	9	US-10-098-263B-3120	Sequence 3120, Ap
c	25	11.2	62.2	25	9	US-10-098-263B-12236	Sequence 12236, A
c	26	11.2	62.2	25	9	US-10-098-263B-26400	Sequence 26400, A
c	27	11.2	62.2	25	9	US-10-098-263B-100906	Sequence 100906,
c	28	11.2	62.2	25	9	US-10-098-263B-120820	Sequence 120820,
c	29	11.2	62.2	25	9	US-10-098-263B-126689	Sequence 126689,
c	30	11.2	62.2	29	9	US-10-062-458-23	Sequence 23, Appl
c	31	11.2	62.2	30	12	US-10-053-632-11	Sequence 11, Appl
c	32	11.2	62.2	30	12	US-10-052-417-11	Sequence 11, Appl
c	33	11.2	62.2	40	10	US-09-245-802-75	Sequence 75, Appl
c	34	11.2	62.2	43	10	US-09-728-574-1	Sequence 1, Appli
c	35	11.2	62.2	44	9	US-10-062-458-18	Sequence 18, Appl
c	36	11.2	62.2	50	9	US-09-741-179A-13	Sequence 13, Appl
c	37	11.2	62.2	50	9	US-09-741-179A-15	Sequence 15, Appl
c	38	11.1	61.1	25	9	US-10-098-263B-26098	Sequence 26098, A
c	39	11.1	61.1	32	10	US-09-919-831-4	Sequence 4, Appli
c	40	10.8	60.0	21	9	US-09-995-529-223	Sequence 223, App
c	41	10.8	60.0	22	10	US-09-815-656-61	Sequence 61, Appl
c	42	10.8	60.0	22	10	US-09-815-656-63	Sequence 63, Appl
c	43	10.8	60.0	22	10	US-09-529-063-114	Sequence 114, App
c	44	10.8	60.0	24	9	US-09-940-185-1700	Sequence 1700, Ap
c	45	10.8	60.0	24	9	US-09-940-185-3740	Sequence 3740, Ap

ALIGNMENTS

RESULT 1
US-09-780-929-98
; Sequence 98, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 98
; LENGTH: 18
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-929-98

Query Match 100.0%; Score 18; DB 10; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.1;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAUGCCUUAUCGGUCGA 18
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DB 1 AAUGCCUUAUCGGUCGA 18

RESULT 2
US-09-780-929-107
; Sequence 107, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity

US-09-940-185-3899

Query Match 65.6%; Score 11.8; DB 9; Length 24;
Best Local Similarity 73.3%; Pred. NO. 3.3e+03;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 4 GGCCUACUGGCGCA 18
||||| ||:||||
DB 2 GGCTAGAGTGCGA 16

RESULT 7

US-09-005-243-36/c
; Sequence 36, Application US/09005243
; Patent No. US20020018763A1
; GENERAL INFORMATION:
; APPLICANT: Zsebo, Krisztina M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Suggs, Sidney V.
; APPLICANT: Martin, Francis H.
; TITLE OF INVENTION: Stem Cell Factor
; NUMBER OF SEQUENCES: 104
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: United States of America
; ZIP: 60606-6402
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/005,243
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/449,653
; FILING DATE: 24-MAY-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/982,255
; FILING DATE: 25-NOV-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/589,701
; FILING DATE: 01-OCT-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/573,616
; FILING DATE: 24-AUG-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/537,198
; FILING DATE: 11-JUN-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/422,383
; FILING DATE: 16-OCT-1989
; ATTORNEY/AGENT INFORMATION:
; NAME: Clough, David W.
; REGISTRATION NUMBER: 36,107
; REFERENCE/DOCKET NUMBER: 01017/34465
; TELEPHONE: 312/474-6300
; TELEFAX: 312/474-0448
; TELEX: 25-3856
; INFORMATION FOR SEQ ID NO: 36:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA

Query Match 64.4%; Score 11.6; DB 10; Length 30;
Best Local Similarity 66.7%; Pred. NO. 4.3e+03;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUACUGGCGCA 18
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DB 19 AAAGGCCTATCCGGCGA 2

RESULT 8

US-09-224-683-36/c
; Sequence 36, Application US/09224683
; Patent No. US20020031491A1
; GENERAL INFORMATION:
; APPLICANT: Zsebo, Krisztina M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Suggs, Sidney V.
; APPLICANT: Martin, Francis H.
; TITLE OF INVENTION: Stem Cell Factor: Composition Claims
; NUMBER OF SEQUENCES: 104
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: United States of America
; ZIP: 60606-6402
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/224,683
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/005,893
; FILING DATE: 12-JAN-1998
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/449,653
; FILING DATE: 24-MAY-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/982,255
; FILING DATE: 25-NOV-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/589,701
; FILING DATE: 01-OCT-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/573,616
; FILING DATE: 24-AUG-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/537,198
; FILING DATE: 11-JUN-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/422,383
; FILING DATE: 16-OCT-1989
; ATTORNEY/AGENT INFORMATION:
; NAME: Clough, David W.
; REGISTRATION NUMBER: 36,107
; REFERENCE/DOCKET NUMBER: 01017/35136
; TELEPHONE: 312/474-6300
; TELEFAX: 312/474-0448
; TELEX: 25-3856
; INFORMATION FOR SEQ ID NO: 36:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single

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; TOPOLOGY: linear
; MOLECULE TYPE: DNA
US-09-224-683-36

Query Match          64.4%; Score 11.6; DB 10; Length 30;
Best Local Similarity 66.7%; Pred. No. 4.3e+03;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGGCGCA 18
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Db 19 AAGGCCCTATCCGGCGCA 2

RESULT 9
US-09-880-508-15/c
; Sequence 15, Application US/09880508
; Publication No. US20030027130A1
; GENERAL INFORMATION:
; APPLICANT: Rice, Charles M.
; Kolykhalov, Alexander A.
; TITLE OF INVENTION: NOVEL 3' TERMINAL SEQUENCE OF HEPATITIS
; C VIRUS GENOME AND DIAGNOSTIC AND THERAPEUTIC USES THEREOF
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Howell & Haferkamp, L.C.
; STREET: 7733 Forsyth Blvd., Suite 1400
; CITY: St. Louis
; STATE: MO
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/880,508
; FILING DATE: 13-Jun-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/897,126
; FILING DATE: <Unknown>
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Henderson, Melodie W.
; REGISTRATION NUMBER: 37,848
; REFERENCE/DOCKET NUMBER: 6029-6836
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 314-727-5188
; TELEFAX: 314-727-6092
; TELEX: <Unknown>
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 46 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 46
; OTHER INFORMATION: /product= "NUCLEOTIDE REPEAT"
; SEQUENCE DESCRIPTION: SEQ ID NO: 15:
US-09-880-508-15

Query Match          64.4%; Score 11.6; DB 9; Length 46;
Best Local Similarity 61.1%; Pred. No. 4.4e+03;
Matches 11; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

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; Sequence 10, Application US/09741179A
; Publication No. US2003002164A1
; GENERAL INFORMATION:
; APPLICANT: MILLS, ALLEN
; TITLE OF INVENTION: DNA-BASED ANALOG NEURAL NETWORKS
; FILE REFERENCE: 31860-168252
; CURRENT APPLICATION NUMBER: US/09/741.179A
; CURRENT FILING DATE: 2000-12-21
; PRIOR APPLICATION NUMBER: 09/129,958

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Db 23 AATGCCCTATTGG 35

Search completed: July 6, 2003, 16:52:33
Job time : 103 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:40:47 ; Search time 1951.09 Seconds
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231.955 Million cell updates/sec

Title: US-09-780-929-98

Perfect score: 18

Sequence: 1 aaagcccaucgugcgca 18

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	18	100.0	29	30	US-09-780-929-107
c 3	13.8	76.7	30	1	PCT-US01-06795A-7
c 5	13.8	76.7	30	1	PCT-US01-06795A-8
6	13.8	76.7	30	30	US-09-798-675-7
7	13.4	74.4	30	30	US-09-798-675-8
8	13.4	74.4	60	34	US-09-908-975-20491
9	13.4	74.4	60	34	US-09-908-975A-20491
10	13.2	73.3	25	36	US-09-956-604-114271
c 11	13.2	73.3	25	36	US-09-956-604-114271
c 12	13.2	73.3	25	36	US-09-956-604-114271
c 13	13.2	73.3	25	36	US-09-956-604-114271
c 14	13.2	73.3	25	36	US-09-956-604-114271
c 15	13.2	73.3	25	36	US-09-956-604-114271
c 16	13.2	73.3	25	36	US-09-956-604-114271
c 17	13.2	73.3	25	36	US-09-956-604-114271
c 18	13.2	73.3	25	36	US-09-956-604B-6465
c 19	13.2	73.3	25	36	US-09-956-604B-105593
c 20	13.2	73.3	25	36	US-09-956-604B-114271
21	13.2	73.3	25	67	US-60-233-166-30098
					Sequence 98, Appl
					Sequence 107, Appl
					Sequence 7, Appl
					Sequence 8, Appl
					Sequence 7, Appl
					Sequence 8, Appl
					Sequence 20491, A
					Sequence 20491, A
					Sequence 20491, A
					Sequence 30098, A
					Sequence 426597, A
					Sequence 6465, Ap
					Sequence 105593, A
					Sequence 114271, A
					Sequence 105593, A
					Sequence 105593, A
					Sequence 114271, A
					Sequence 6465, Ap
					Sequence 105593, A
					Sequence 114271, A
					Sequence 30098, A

PCT-US01-06795A-8

Query Match 76.7%; Score 13.8; DB 1; Length 30;
Best Local Similarity 64.7%; Pred. No. 2.4e+03;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUACGUGCGGA 18
I:||||:|:|:|:|:|:|
Db 6 ATGGCGTATCGATCGGA 22

RESULT 5

US-09-798-675-7/c
; Sequence 7, Application US/09798675
; GENERAL INFORMATION:
; APPLICANT: Emory University
; TITLE OF INVENTION: HIV VACCINES
; FILE REFERENCE: E056 2020
; CURRENT APPLICATION NUMBER: US/09/798,675
; CURRENT FILING DATE: 2001-12-11
; PRIOR APPLICATION NUMBER: US 60/186,364
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/251,083
; PRIOR FILING DATE: 2000-12-01
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 7
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:

US-09-798-675-7
OTHER INFORMATION: primer for site-directed mutagenesis for introducing Cla I site

Query Match 76.7%; Score 13.8; DB 30; Length 30;
Best Local Similarity 64.7%; Pred. No. 2.4e+03;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUACGUGCGGA 18
I:||||:|:|:|:|:|:|
Db 25 ATGGCGTATCGATCGGA 9

RESULT 6

US-09-798-675-8
; Sequence 8, Application US/09798675
; GENERAL INFORMATION:
; APPLICANT: Emory University
; TITLE OF INVENTION: HIV VACCINES
; FILE REFERENCE: E056 2020
; CURRENT APPLICATION NUMBER: US/09/798,675
; CURRENT FILING DATE: 2001-12-11
; PRIOR APPLICATION NUMBER: US 60/186,364
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/251,083
; PRIOR FILING DATE: 2000-12-01
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:

US-09-798-675-8
OTHER INFORMATION: primer for site-directed mutagenesis to introduce Cla I site

Query Match 76.7%; Score 13.8; DB 30; Length 30;
Best Local Similarity 64.7%; Pred. No. 2.4e+03;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUACGUGCGGA 18
I:||||:|:|:|:|:|:|
Db 6 ATGGCGTATCGATCGGA 22

RESULT 7
US-09-908-975-20491
; Sequence 20491, Application US/09908975
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: MINTZ, Liat
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SI
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME
; FILE REFERENCE: 36688-0005
; CURRENT APPLICATION NUMBER: US/09/908,975
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20491
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-908-975-20491

Query Match 74.4%; Score 13.4; DB 34; Length 60;
Best Local Similarity 66.7%; Pred. No. 4.4e+03;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 AAUGGCCUACGUGG 15
I:||||:|:|:|:|:|:|
Db 1 ATGGCCTATCGGTG 15

RESULT 8

US-09-908-975A-20491
; Sequence 20491, Application US/09908975A
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: MINTZ, Liat
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SI
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME
; FILE REFERENCE: 36688-0006
; CURRENT APPLICATION NUMBER: US/09/908,975A
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20491
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-908-975A-20491

Query Match 74.4%; Score 13.4; DB 34; Length 60;
Best Local Similarity 66.7%; Pred. No. 4.4e+03;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 AAUGGCCUACGUGG 15
I:||||:|:|:|:|:|:|
Db 1 ATGGCCTATCGGTG 15

RESULT 9

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US-60-287-724-20491
; Sequence 20491, Application US/60287724
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICING
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME
; FILE REFERENCE: 36688-0004
; CURRENT APPLICATION NUMBER: US/60/287,724
; CURRENT FILING DATE: 2001-05-02
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20491
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
US-60-287-724-20491

Query Match          74.4%; Score 13.4; DB 72; Length 60;
Best Local Similarity 66.7%; Pred. No. 4.4e+03;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUG 15
| :|||:|:|:|:|
Db 1 ATTGGCCTATCGGTG 15

RESULT 10
US-09-954-427-30098
; Sequence 30098, Application US/09954427
; GENERAL INFORMATION:
; APPLICANT: Mitmann
; APPLICANT: Affymetrix, Inc.
; TITLE OF INVENTION: Methods of Genetic Analysis of the Rat
; TITLE OF INVENTION: Genome
; FILE REFERENCE: 3112
; CURRENT APPLICATION NUMBER: US/09/954,427
; CURRENT FILING DATE: 2001-09-17
; NUMBER OF SEQ ID NOS: 420907
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 30098
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; PUBLICATION INFORMATION:
; DATABASE ACCESSION NUMBER: GenBank AA818650
US-09-954-427-30098

Query Match          73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 66.7%; Pred. No. 5.3e+03;
Matches 12; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCA 18
|:|:|:|:|:|:|
Db 8 AATGCTCTATCGCGCAA 25

RESULT 11
US-09-956-584-426597/c
; Sequence 426597, Application US/09956584
; GENERAL INFORMATION:
; APPLICANT: Mitmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Mus Musculus
; FILE REFERENCE: 3115.1
; CURRENT APPLICATION NUMBER: US/09/956,584
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,017
; PRIOR FILING DATE: 2000-09-20
; NUMBER OF SEQ ID NOS: 605887
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1

US-09-956-604-6465/c
; Sequence 6465, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mitmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 105593
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-105593

Query Match          73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18
|:|:|:|:|:|:|
Db 20 AGTGGCCTATCAGTGCCA 3

RESULT 14
US-09-956-604-114271/c
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; SEQ ID NO 426597
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-956-584-426597

Query Match          73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 66.7%; Pred. No. 5.3e+03;
Matches 12; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18
|:|:|:|:|:|:|
Db 21 AAGGACCTATCGGTCCGA 4

RESULT 12
US-09-956-604-6465/c
; Sequence 6465, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mitmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 6465
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-6465

Query Match          73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18
|:|:|:|:|:|:|
Db 23 AATGGTCTATCGTTGAGA 6

RESULT 13
US-09-956-604-105593/c
; Sequence 105593, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mitmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 105593
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-105593

Query Match          73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18
|:|:|:|:|:|:|
Db 23 AATGGTCTATCGTTGAGA 6

RESULT 13
US-09-956-604-105593/c
; Sequence 105593, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mitmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 105593
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-105593

Query Match          73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18
|:|:|:|:|:|:|
Db 20 AGTGGCCTATCAGTGCCA 3

RESULT 14
US-09-956-604-114271/c
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; Sequence 114271, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956.604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234, 049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 114271
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-114271

Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY      1 AAUGGCCUAUCGUGCGGA 18
DB      20 AGTGGCTATCGCTACGA 3

RESULT 15
US-09-956-604A-6465/c
; Sequence 6465, Application US/09956604A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956.604A
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234, 049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 6465
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604A-6465

Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY      1 AAUGGCCUAUCGUGCGGA 18
DB      23 AATGGTCTATCGTTGAGA 6

Search completed: July 6, 2003, 16:29:55
Job time : 1953.09 secs
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	Matches	11;	Conservative	4;	Mismatches	2;	Indels	0;	Gaps	0;
Qy	2	AUGGCCUACGUGCGCA	18							
		: : : : : : : :								
Db	25	ATGCCGTATCGATCCGA	9							

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RESULT 2
US-10-336-566-22
; Sequence 22, Application US/10336566
; GENERAL INFORMATION:
; APPLICANT: Robinson, Harriet L.
; APPLICANT: Smith, James M.
; APPLICANT: Hua, Jian
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR GENERATING
; TITLE OF INVENTION: AN IMMUNE RESPONSE
; FILE REFERENCE: 12804-006001
; CURRENT APPLICATION NUMBER: US/10/336,566
; CURRENT FILING DATE: 2003-01-03
; PRIOR APPLICATION NUMBER: US 10/093,953
; PRIOR FILING DATE: 2002-03-08
; PRIOR APPLICATION NUMBER: US 09/798,675
; PRIOR FILING DATE: 2001-03-01
; PRIOR APPLICATION NUMBER: PCT/US01/06795
; PRIOR FILING DATE: 2001-03-02
; PRIOR APPLICATION NUMBER: US 60/251,083
; PRIOR FILING DATE: 2000-12-01
; PRIOR APPLICATION NUMBER: US 60/186,364
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/324,845
; PRIOR FILING DATE: 2001-09-25
; NUMBER OF SEQ ID NOS: 87
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 22
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Primer
US-10-336-566-22

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Query Match	76.78;	Score 13.8;	DB 15;	Length 30;
Best Local Similarity	64.78;	Pred. No. 9.6+0.2;		
Matches 11;	Conservative 4;	Mismatches 2;	Indels 0;	Gaps 0;
Qy	2	AUGGCCUAUGGGGCGA	18	
Db	6	ATGGCGTATCGATCGCA	22	

RESULT 3
 US-10-310-188-56966/c
 : Sequence 56966, Application US/10310188
 : GENERAL INFORMATION:
 : APPLICANT: RosettaGenomics
 : TITLE OF INVENTION: BIOINFORMATICALLY DETECTABLE GROUP OF NOVEL VIRAL REGULATORY GENES
 : TITLE OF INVENTION: US/10310188
 : FILE REFERENCE: 47487
 : CURRENT APPLICATION NUMBER: US/10/310,188
 : CURRENT FILING DATE: 2002-12-19
 : NUMBER OF SEQ ID NOS: 86841
 : SOFTWARE: PatentIn version 3.1
 : SEQ ID NO 56966
 : LENGTH: 24
 : TYPE: DNA
 : ORGANISM: Homo sapiens
 US-10-310-188-56966

```

Query Match          73.3%; Score 13.2; DB 14; Length 24;
Best Local Similarity 61.1%; Pred. No. 2.1e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;
Qv      1 AAUGGCCUACGUGCGCA 18

```

```

      ||:|||||:|:|:| |
Db      21  AATGGCCTATGGATGCAA  4

RESULT 4
US-60-427-808-19526
; Sequence 19526, Application US/60427808
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Mouse
; FILE REFERENCE: 3528
; CURRENT APPLICATION NUMBER: US/60/427,808
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 982914
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 19526
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Mus musculus
US-60-427-808-19526

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```

Query Match      73.3%  Score 13.2; DB 18; Length 25;
Best Local Similarity 61.1%  Pred. No. 2.1e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGGCA 18
   ||:|:|:|:|:|
Db 2 AATAGCTATCGTTGGCA 19

```

```

RESULT 5
US-60-427-808-125229/c
; Sequence 125229, Application US/60427808
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Mouse
; FILE REFERENCE: 3528
; CURRENT APPLICATION NUMBER: US/60/427,808
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 982914
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 125229
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Mus musculus
US-60-427-808-125229.

```

Query Match	73.3%	Score 13.2;	DB 18;	Length 25;
Best Local Similarity	66.7%	Pred. No. 2.le+03;		
Matches 12;	Conservative 3;	Mismatches 3;	Indels 0;	Gaps 0;
Qy	1	AAUGGCCUAUCGGGCGA	18	
		: :		
Db	22	AAGGCCATCAGTACGA	5	
		: :		

```

RESULT 6
US-60-427-808-897906
; Sequence 897906, Application US/60427808
; GENERAL INFORMATION:
; APPLICANT: xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Mouse
; FILE REFERENCE: 3528
; CURRENT APPLICATION NUMBER: US/60/427,808
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 982914
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 897906
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Mus musculus
US-60-427-808-897906

```

0v 6 CCUAUCGGUGCGA 18

Db 11:11:11:1111
2 CCTATCGGTGCGA 14

RESULT 12

US-60-427-808-512279/c
; Sequence 512279, Application US/60427808
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Mouse
; FILE REFERENCE: 3528
; CURRENT APPLICATION NUMBER: US/60/427,808
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 982914
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 512279
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Mus musculus
US-60-427-808-512279

Query Match 72.2%; Score 13; DB 18; Length 25;
Best Local Similarity 76.9%; Pred. No. 2.7e+03;
Matches 10; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGG 13
11:1111:1:1111
Db 23 AATGGCCTATCGG 11

RESULT 13

US-09-954-445A-39955/c
; Sequence 39955, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954,445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 39955
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-39955

Query Match 71.1%; Score 12.8; DB 11; Length 25;
Best Local Similarity 62.5%; Pred. No. 3.5e+03;
Matches 10; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGC 16
11:11:11:1111
Db 22 AATGCATATCGGTGC 7

RESULT 14

US-10-355-577-133076
; Sequence 133076, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-UI133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 133076
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien

US-10-355-577-133076

Query Match 71.1%; Score 12.8; DB 14; Length 25;
Best Local Similarity 68.8%; Pred. No. 3.5e+03;
Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
QY 2 AUGGCCUAUCGGUGCG 17
1111:1111:1111
Db 5 AAGGACTATCGGTGCG 20

RESULT 15

US-10-355-577-152466/c
; Sequence 152466, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-UI133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 152466
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-152466

Query Match 71.1%; Score 12.8; DB 14; Length 25;
Best Local Similarity 62.5%; Pred. No. 3.5e+03;
Matches 10; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 3 UGGCCUAUCGGUGCGA 18
11:11:11:1111
Db 21 TTGACTATCGGTGCGA 6

Search completed: July 6, 2003, 16:49:14
Job time : 625.545 secs